

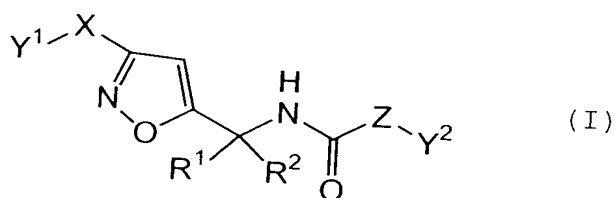
## DESCRIPTION

SUBSTITUTED ISOXAZOLE ALKYLAMINE DERIVATIVE AND AGRI-  
HORTICULTURAL FUNGICIDE

## 5 TECHNICAL FIELD

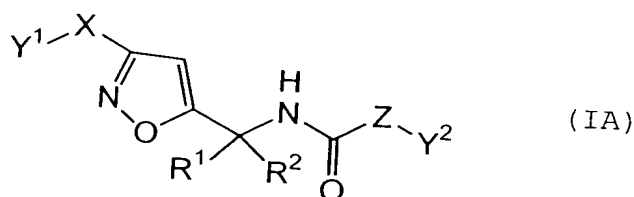
The present invention relates to:

- 1) a novel substituted isoxazole derivative represented by the formula (I):



- 10 (symbols in the formula are as described later); and

- 2) an agri-horticultural fungicide containing a substituted isoxazole derivative represented by the formula (IA):



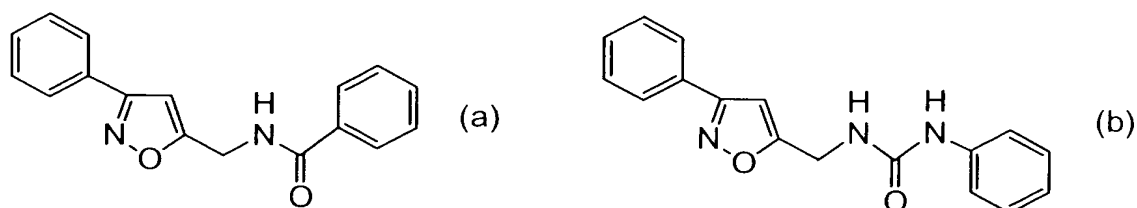
- 15 (symbols in the formula are as described later).

## BACKGROUND ART

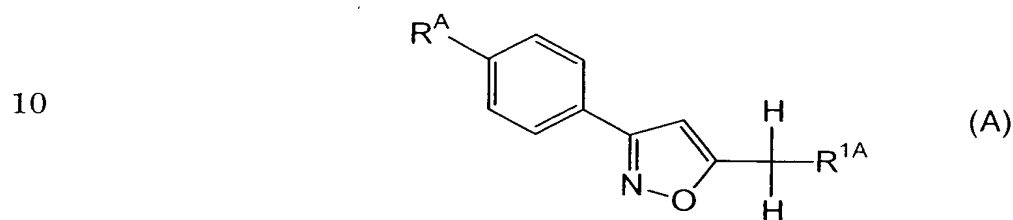
- 1) JP 9-509951 A, International Publication WO 98/28282, US Patent No. 5,859,257, British Patent No. 1,067,203, and European Patent No. 277,791 disclose compounds of substituted isoxazole alkylamine derivatives,

but have no description about fungicidal activity.

2) In J. C. S. Perkin Trans. I, 16, 1694-1696 (1976),  
a spirooxazoline ring formation reaction is performed using  
the following compounds (a) and (b) as starting materials,  
5 but there is no description about fungicidal activity of the  
compounds (a) and (b).



3) JP 52-154525 A shows that a compound represented by  
the following general formula (A):



(wherein  $R^A$  represents a chloro, bromo, fluoro, or  
trifluoromethyl group or a hydrogen atom, and  $R^{1A}$  represents  
a bromo, chloro, isothiocyanate, or amino group, amino-  
hydrochloride or hydrobromide,  $-NHCSNH(C1 \text{ to } C3)\text{alkyl}$ ,  $-$   
15  $NHCOO(C1 \text{ to } C3)\text{alkyl}$ , or 3,5,7-triaza-1-azonia adamantyl  
chloride or bromide) serves as fungicide to exhibit disease  
control activity particularly against tomato late blight,  
bean powdery mildew, cucumber anthracnose, rice blast, wheat  
leaf spot, grape gray mold, apple scab, grape downy mildew,

and sugar beet leaf spot. However, the activity as the fungicide of the compound described in the specification of this patent document is not sufficient.

In cultivation of agricultural or horticultural crops, various disease damages which seriously affect productive efficiency have been known, and examples of the vital disease damages include: rice blast and sheath blight in rice cultivation; and powdery mildew, downy mildew, and gray mold in cultivation of fruits, vegetables, and grain crops.

Various fungicides have been used against the disease damages caused by those plant pathogenic fungi. However, some fungicides known publicly and used conventionally are not effective enough any longer because fungi resistant to each of the fungicides occur owing to long-continued use of the fungicides, or some of the fungicides can exhibit sufficient fungicidal effects only when used in combination with other drugs. Therefore, in the industrial field of agricultural or horticultural cultivation, a novel compound for preventing/treating plant disease damages, with a chemical skeleton different from conventional drugs, which exerts a sufficient preventive effect, has been long demanded. The invention is to solve problems involved in the prior art mentioned above. An object of the invention is to provide a novel drug which, owing to its chemical skeleton different from those of conventionally known drugs

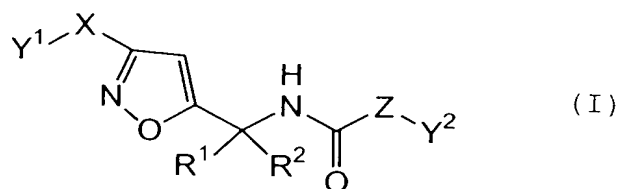
exerts a sufficient activity for preventing plant disease damage without cross-reacting with resistant fungi.

#### DISCLOSURE OF THE INVENTION

5           In order to accomplish the above object, the inventors of the present invention have synthesized novel substituted isoxazole alkylamine derivatives whose activity for preventing plant disease damages has not been known, and they have intensively studied on the activity for preventing  
10 plant disease damages and usefulness of the derivatives. As a result, the inventors have found that a substituted isoxazole alkylamine derivative represented by the formula (I) or the formula (IA) described below has an excellent plant pathogenic fungus control effect without giving a  
15 harmful effect to plants.

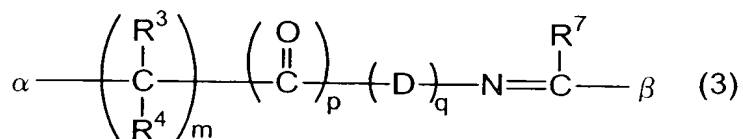
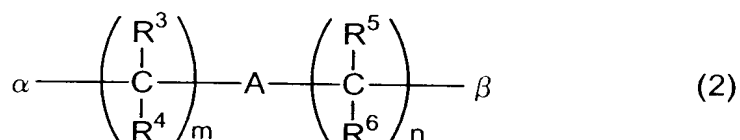
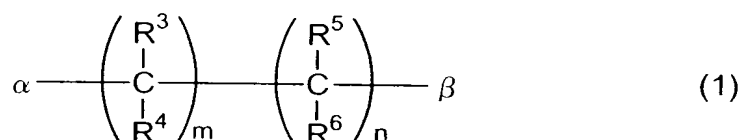
That is, the present invention relates to an agent for preventing plant diseases which is for use in the agricultural field, comprising as active ingredient a substituted isoxazole alkylamine derivative represented by  
20 the following general formula (I) or a substituted isoxazole alkylamine derivative represented by the following general formula (IA).

A substituted isoxazole alkylamine derivative denoted by the formula (I):



wherein  $R^1$  and  $R^2$  may be the same or different, and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkenyl group, a lower alkynyl group, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxy carbonyl group, a lower alkylthio group, a halogen atom, a hydroxyl group, a carboxyl group, or a cyano group, or  $R^1$  and  $R^2$  may together form a cycloalkyl group which may be substituted;

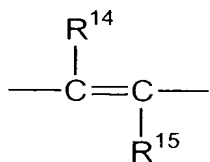
X represents the following formula (1), (2), or (3):



(wherein  $R^3$ ,  $R^4$ ,  $R^5$ , and  $R^6$  may be the same or different, and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkenyl group, a lower alkynyl group, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxy carbonyl group, a lower alkylthio group, a halogen atom, a hydroxyl group, a carboxyl group, or a cyano group, or  $R^3$  and  $R^4$  may together form a cycloalkyl group which may be substituted;

alkylthio group, an amino group, a lower alkylamino group, a di(lower alkyl)amino group, a halogen atom, a hydroxyl group, or a cyano group, or each pair of R<sup>3</sup> and R<sup>4</sup>, and R<sup>5</sup> and R<sup>6</sup> may together form a cycloalkyl group which may be substituted;

A represents an oxygen atom, a sulfur atom, an -S(O)- group, an -S(O)<sub>2</sub>- group, an -NR<sup>12</sup>- group (wherein R<sup>12</sup> represents a hydrogen atom or a lower alkyl group), a carbonyl group, an -NH-CO- group, a -CO-NH- group, a -C≡C- group, an -NH-CO-NH- group, an -O-CONH- group, an -HC=N- group or a group represented by the formula below



(wherein R<sup>14</sup> and R<sup>15</sup> each represent a hydrogen atom or a lower alkyl group);

m and n each represent 0 or an integer of 1 to 3;

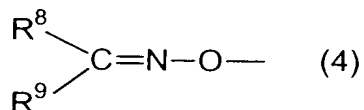
p and q each represent 0 or 1;

R<sup>7</sup> represents a hydrogen atom or a lower alkyl group; and

D represents an oxygen atom or an -NH- group, wherein α binds to a Y<sup>1</sup> side, and β binds to an isoxazole moiety);

Y<sup>1</sup> represents a lower alkyl group which may be substituted, a lower alkenyl group which may be substituted, a lower alkynyl group which may be substituted, a cycloalkyl group which may be substituted, a lower cyclo alkenyl group

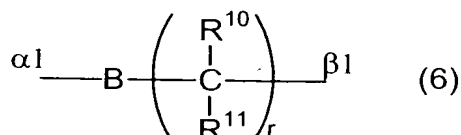
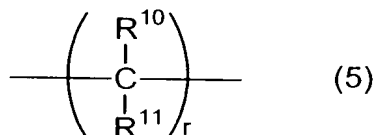
which may be substituted, a phenyl group which may be substituted, a naphthyl group which may be substituted, a heteroaryl group which may be substituted, an aliphatic hetero ring which may be substituted, or the following  
 5 formula (4):



(wherein  $\text{R}^8$  and  $\text{R}^9$  each represent a hydrogen atom, a lower alkyl group which may be substituted, or a phenyl group which may be substituted, or  $\text{R}^8$  and  $\text{R}^9$  may together form a  
 10 cycloalkyl group which may be substituted);

$\text{Y}^2$  represents a lower alkenyl group which may be substituted, a lower alkynyl group which may be substituted, a cycloalkyl group which may be substituted, a lower cyclo alkenyl group which may be substituted, a phenyl group which  
 15 may be substituted, a naphthyl group which may be substituted, a heteroaryl group which may be substituted, or an aliphatic hetero ring which may be substituted;

$\text{Z}$  represents a group denoted by the following formula (5) or (6):



(wherein  $\text{R}^{10}$  and  $\text{R}^{11}$  may be the same or different, and each represent a hydrogen atom, a lower alkyl group which may be

substituted, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxycarbonyl group, a lower alkylthio group, an amino group, a lower alkylamino group, a di(lower alkyl)amino group, a halogen atom, a hydroxyl group, or a cyano group, or  $R^{10}$  and  $R^{11}$  may together form a cycloalkyl group which may be substituted;

$r$  represents 0 or an integer of 1 to 3; and

$B$  represents an oxygen atom, a sulfur atom, or an  $-NR^{13}-$  group wherein  $R^{13}$  represents a hydrogen atom or a lower alkyl group), wherein any one of  $\alpha 1$  and  $\beta 1$  may be bound to a  $Y^2$  side;

provided that the following (1) to (3) are excluded:

(1) a compound in which  $X$  is a single bond (i.e. the case that each of  $m$  and  $n$  is 0 in the formula (1)) and  $Y^1$  represents a 4-hydroxy-3,5-di-tert-butylphenyl group;

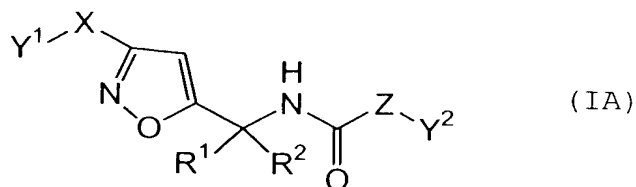
(2) a compound in which  $R^1$  and  $R^2$  represent hydrogen atoms,  $X$  is a single bond (i.e. the case that each of  $m$  and  $n$  is 0 in the formula (1)), and both  $Y^1$  and  $Y^2$  represent unsubstituted phenyl groups (2-1) when  $Z$  is a single bond (i.e. the case that  $r$  is 0 in the formula (5)) or (2-2) when  $Z$  is an NH group (i.e. the case that  $r=0$  and  $B$  represents an  $-NH-$  group in the formula (6)); and

(3) a compound in which  $R^1$  and  $R^2$  represent hydrogen atoms,  $Z$  is a single bond (i.e. the case that  $r$  is 0 in the formula (5)), and  $Y^2$  represents a 4-hydroxycinnolin-3-yl group which



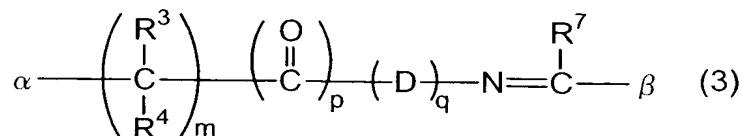
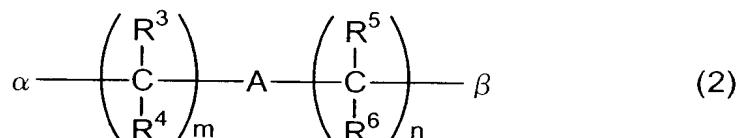
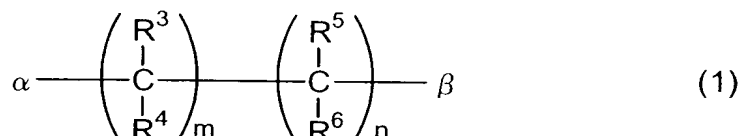
may be substituted.

A substituted isoxazole alkylamine derivative represented by the formula (IA):



5 wherein R<sup>1</sup> and R<sup>2</sup> may be the same or different, and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkenyl group, a lower alkynyl group, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxycarbonyl group, a lower alkylthio group,  
 10 a halogen atom, a hydroxyl group, a carboxyl group or a cyano group, or R<sup>1</sup> and R<sup>2</sup> may together form a cycloalkyl group which may be substituted;

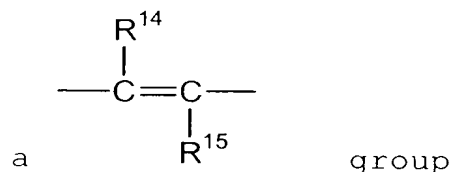
X represents the following formula (1), (2) or (3):



15 (wherein R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup>, and R<sup>6</sup> may be the same or different, and each represent a hydrogen atom, a lower alkyl group which

may be substituted, a lower alkenyl group, a lower alkynyl group, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxy carbonyl group, a lower alkylthio group, an amino group, a lower alkylamino group, a di(lower alkyl)amino group, a halogen atom, a hydroxyl group, or a cyano group, or each pair of  $R^3$  and  $R^4$ , and  $R^5$  and  $R^6$  may together form a cycloalkyl group which may be substituted;

A represents an oxygen atom, a sulfur atom, an  $-S(O)-$  group, an  $-S(O)_2-$  group, an  $-NR^{12}-$  group (wherein  $R^{12}$  represents a hydrogen atom or a lower alkyl group), a carbonyl group, an  $-NH-CO-$  group, a  $-CO-NH-$  group, a  $-C\equiv C-$  group, an  $-NH-CO-NH-$  group, an  $-O-CONH-$  group, an  $-HC=N-$  group, or



(wherein  $R^{14}$  and  $R^{15}$  each represent a hydrogen atom or a lower alkyl group);

m and n each represent 0 or an integer of 1 to 3;

p and q each represent 0 or 1;

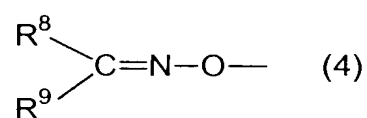
$R^7$  represents a hydrogen atom or a lower alkyl group; and

D represents an oxygen atom or an  $-NH-$  group,

a binds to a  $Y^1$  side, and  $\beta$  binds to an isoxazole moiety);

$Y^1$  represents a lower alkyl group which may be substituted, a lower alkenyl group which may be substituted,

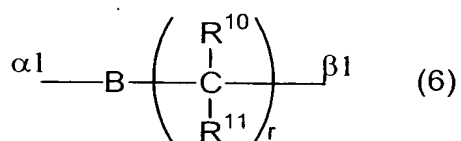
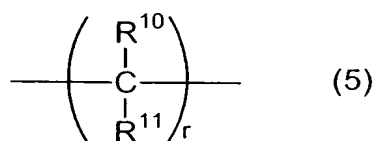
a lower alkynyl group which may be substituted, a cycloalkyl group which may be substituted, a lower cyclo alkenyl group which may be substituted, a phenyl group which may be substituted, a naphthyl group which may be substituted, a  
 5 heteroaryl group which may be substituted, an aliphatic hetero ring which may be substituted, or the following formula (4):



(wherein  $\text{R}^8$  and  $\text{R}^9$  each represent a hydrogen atom, a lower  
 10 alkyl group which may be substituted, or a phenyl group which may be substituted, or  $\text{R}^8$  and  $\text{R}^9$  may together form a cycloalkyl group which may be substituted);

$\text{Y}^2$  represents a lower alkenyl group which may be substituted, a lower alkynyl group which may be substituted,  
 15 a cycloalkyl group which may be substituted, a lower cyclo alkenyl group which may be substituted, a phenyl group which may be substituted, a naphthyl group which may be substituted, a heteroaryl group which may be substituted, or an aliphatic hetero ring which may be substituted;

20  $\text{Z}$  represents a group represented by the following formula (5) or (6):



(wherein  $R^{10}$  and  $R^{11}$  may be the same or different, and each represent a hydrogen atom, a lower alkyl group which may be substituted, a cycloalkyl group which may be substituted, a lower alkoxy group, a lower alkoxycarbonyl group, a lower alkylthio group, an amino group, a lower alkylamino group, a di(lower alkyl)amino group, a halogen atom, a hydroxyl group, or a cyano group, or  $R^{10}$  and  $R^{11}$  may together form a cycloalkyl group which may be substituted;

$r$  represents 0 or an integer of 1 to 3; and

10  $B$  represents an oxygen atom, a sulfur atom, or an  $-NR^{13}-$  group wherein  $R^{13}$  represents a hydrogen atom or a lower alkyl group), wherein any one of  $\alpha 1$  and  $\beta 1$  may be bound to a  $Y^2$  side.

[Compounds of the Present Invention]

15 In the formula (I) or (IA), lower alkyl group refers to an alkyl group having 1 to 6 carbon atoms such as methyl, ethyl, propyl, butyl, pentyl, and hexyl groups, and isomer groups thereof.

20 The lower alkenyl group refers to an alkenyl group having 1 to 3 double bonds and 2 to 6 carbon atoms such as vinyl, propenyl, butenyl, butadienyl, and hexatrienyl groups, and isomer groups thereof.

25 The lower alkynyl group refers to an alkynyl group having 1 to 3 triple bonds and 2 to 6 carbon atoms such as an ethynyl, propynyl, pentynyl, and hexynyl group, and

isomer groups thereof.

The cycloalkyl group refers to a cycloalkyl group having 3 to 8 carbon atoms such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, and cyclooctyl groups.

5       The cycloalkenyl group refers to a cycloalkenyl group having 3 to 8 carbon atoms such as cyclopropynyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, cycloheptenyl, and cyclooctenyl groups, and isomer groups thereof.

10       The lower alkoxy group refers to an alkoxy group having 1 to 6 carbon atoms such as a methyloxy group, an ethyloxy group, a propyloxy group, a butyloxy group, a pentyloxy group, a hexyloxy group, and isomer groups thereof.

15       The lower alkoxycarbonyl group refers to an alkoxycarbonyl group having 1 to 6 carbon atoms such as a methyloxycarbonyl group, an ethyloxycarbonyl group, a propyloxycarbonyl group, a butyloxycarbonyl group, a pentyloxycarbonyl group, a hexyloxycarbonyl group, and isomer groups thereof.

20       The lower alkylthio group refers to an alkylthio group having 1 to 6 carbon atoms such as a methylthio group, an ethylthio group, a propylthio group, a butylthio group, a pentylthio group, a hexylthio group, and isomer groups thereof.

25       The halogen atom refers to a fluorine atom, a chlorine atom, a bromine atom, or an iodine atom.

The heteroaryl group refers to an aromatic monocyclic or bicyclic hetero ring having 1 to 3 hetero atoms including an oxygen atom, a sulfur atom, and a nitrogen atom as a ring forming atom. Examples thereof include pyrrole, pyrazole, imidazole, triazole, pyridine, pyrimidine, triazine, furan, thiophene, thiazole, oxazole, isoxazole, isothiazole, benzothiazole, benzoxazole, indazole, indole, quinoline, purine, and pteridine rings.

The aliphatic hetero ring refers to a 5 to 6-membered aliphatic hetero ring having 1 to 3 hetero atoms including an oxygen atom, a sulfur atom, and a nitrogen atom as a ring forming atom. Examples thereof include a pyrrolidine ring, a piperidine ring, a piperazine ring, a tetrahydropyran ring, a tetrahydrofuran ring, a morpholine ring, and dioxane.

A lower alkyl group which may be substituted, a lower alkenyl group which may be substituted, a lower alkynyl group which may be substituted, a cycloalkyl group which may be substituted, a cycloalkenyl group which may be substituted, lower alkoxy group which may be substituted, or an aliphatic hetero ring which may be substituted may be substituted with one or more substituents selected from: a lower alkoxy group; a lower alkoxycarbonyl group; a halogen atom; a cyano group; a nitro group; a phenyl group which may be substituted; a lower acyl group; and a lower acyloxy group.

A phenyl group which may be substituted, a phenyloxy group which may be substituted, a naphthyl group which may be substituted, or a heteroaryl group which may be substituted may be substituted with one or more substituents  
5 selected from: a lower alkyl group which may be substituted with a halogen atom or a lower alkoxy group; a lower alkenyl group which may be substituted; a lower alkynyl group which may be substituted; a lower alkoxy group which may be substituted with a halogen atom and like; a lower alkylthio  
10 group; a lower alkanesulfinyl group; a lower alkanesulfonyl group; an  $R^{16}$ -NHCO- group; an  $R^{16}$ -CONH- group (wherein,  $R^{16}$  represents a hydrogen atom or a lower alkyl group); an  $R^{17}$ -O-NHCO- group (wherein,  $R^{17}$  represents a lower alkyl group); an  $R^{18}$ -CO- group (wherein,  $R^{18}$  represents a hydrocarbon group  
15 (for example, a lower alkyl group and a phenyl group) which may be substituted with a halogen atom and like); a lower alkoxy carbonyl group; a carboxyl group; a cycloalkyl group which may be substituted; an amino group; a lower alkylamino group; a lower dialkylamino group; an  $R^{19}$ O-N=C( $R^{20}$ )- group  
20 (wherein,  $R^{19}$  represents a hydrogen atom or a lower alkyl group, and  $R^{20}$  represents a hydrogen atom, a lower alkyl group, or an amino group); an aliphatic heterocyclo ring group; a lower alkylcarbonylhydrazino group; a lower alkyloxy carbonylhydrazino group; a formyl group; an  
25  $H_2NN=C(R^{21})$ - group; an  $R^{22}(O)CHNN=C(R^{21})$ - group (wherein,  $R^{21}$

and  $R^{22}$  each represent a hydrogen atom or a lower alkyl group); a hydroxyl group; a phenyl group which may be substituted; a phenyloxy group which may be substituted; a naphthyl group which may be substituted; a heteroaryl group  
5 which may be substituted; a halogen atom; a cyano group; and a nitro group. Further, adjacent substituents together may form a cyclic alkyl group which may be substituted or a cyclic ether group which may be substituted.

A preferable X may be provided as follows. In the  
10 above-mentioned formula (1), m and n are each 0 or 1, more specifically  $m = n = 0$  (single bond),  $m + n = 1$  (a methylene group which may be substituted), and  $m = n = 1$  (an ethylene group which may be substituted). In the above-mentioned formula (2), m and n are preferably each 0 or 1, more  
15 preferably  $m = n = 0$  (A),  $m = 1$  and  $n = 0$ , and  $m = 0$  and  $n = 1$ , even more preferably  $m = n = 0$  (A), and  $m = 0$  and  $n = 1$ . In the above-mentioned formula (3), preferable examples of the  $-(CO)_p-(D)_q-$  include a single bond, an  $-O-$  group, and a  $-CONH-$  group.

20 A preferable Z is a group provided in the above-mentioned formulae (5) and (6), where r is 0 or 1. More preferable examples of Z include a single bond, a  $-CH(R^{10a})-$  group (where,  $R^{10a}$  represents a hydrogen atom, a C1 to C3 alkyl group, a halogen atom, or a cyano group), an  $-O-$  group,  
25 an  $-NH-$  group, an  $-N(CH_3)-$  group, an  $-O-C(R^{11a})_2-$  group, a -



$C(R^{11a})_2-O-$  group, and an  $-NH-C(R^{11a})_2-$  group (where,  $R^{11a}$  independently represents a hydrogen atom or C1 to C3 alkyl group).

Preferable examples of  $Y^1$  include: a lower alkyl group  
5 which may be substituted; a cycloalkyl group which may be substituted; a phenyl group which may be substituted; and a heteroaryl group which may be substituted.

Preferable examples of  $Y^2$  include: a cycloalkyl group which may be substituted; a phenyl group which may be substituted;  
10 substituted; a naphthyl group which may be substituted; and a heteroaryl group which may be substituted. More Preferred examples of  $Y^2$  include: a cycloalkyl group which may be substituted; a phenyl group which may be substituted; and a naphthyl group which may be substituted.

15 Examples of the compound represented by the formula (I) or (IA) are preferably those represented below.

[1] A substituted isoxazole alkylamine derivative, in which  $R^1$  and  $R^2$  each independently represent a hydrogen atom or a lower alkyl group which may be substituted.

20 [2] The substituted isoxazole alkylamine derivative according to [1], in which Z is an oxygen atom (when r is 0 and B is an oxygen atom in the formula (6)), and  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl  
25 group which may be substituted.

[3] The substituted isoxazole alkylamine derivative according to [1], in which Z represents an  $-NR^{13}-$  group (wherein  $R^{13}$  represents a hydrogen atom or a lower alkyl group) (when r is 0 and B is an  $-NR^{13}-$  group) and  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted.

[4] The substituted isoxazole alkylamine derivative according to [1], in which X and Z are single bonds (when m and n are 0 in the formula (1) and r is 0 in the formula (5)) and  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted.

[5] The substituted isoxazole alkylamine derivative according to [1], in which X represents a group represented by the formula (1) wherein  $R^3$ ,  $R^4$ ,  $R^5$ , and  $R^6$  may be the same or different and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkoxy group, a lower alkoxycarbonyl group, a halogen atom, or a cyano group,  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted, and  $m + n$  is 1, 2, or 3.

[6] The substituted isoxazole alkylamine derivative according to [1], in which X represents a group represented by the formula (2) wherein  $R^3$ ,  $R^4$ ,  $R^5$ , and  $R^6$  may be the same

or different and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkoxy group, a lower alkoxy carbonyl group, a halogen atom, or a cyano group,  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted,  $n$  is 1 or 2, and  $m + n$  is 1, 2, or 3.

[7] The substituted isoxazole alkylamine derivative according to [1], in which  $X$  represents a group represented by the formula (2) wherein  $R^3$ ,  $R^4$ ,  $R^5$ , and  $R^6$  may be the same or different and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkoxy group, a lower alkoxy carbonyl group, a halogen atom, or a cyano group,  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted,  $n$  is 0, and  $m$  is 0, 1, or 2.

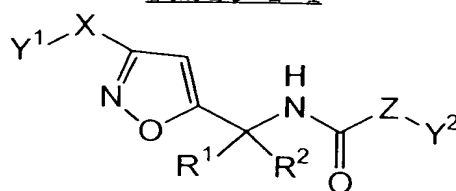
[8] The substituted isoxazole alkylamine derivative according to [1], in which  $X$  represents a group denoted by the formula (3) wherein  $R^3$  and  $R^4$  may be the same or different and each represent a hydrogen atom, a lower alkyl group which may be substituted, a lower alkoxy group, a halogen atom, or a cyano group,  $m$  represents 0, 1, or 2, and  $Y^2$  represents a phenyl group which may be substituted, a naphthyl group which may be substituted, or a cycloalkyl group which may be substituted.

[9] The substituted isoxazole alkylamine derivative according to [1], in which  $Y^1$  represents a methyl group substituted with a halogen atom, and Z is a single bond (when m and n are 0 in the formula (1)).

5        Next, representative examples of the compounds of the invention represented by the formula (I) are shown in Tables 1-1 to 4-8, but the compounds of the invention are not limited thereto (the same number is used as Compound Number throughout the specification.).

10        In the table, Me represents a methyl group, Et represents an ethyl group, n-Pr represents a normal propyl group, i-Pr represents an isopropyl group, n-Bu represents a normal butyl group, i-Bu represents an isobutyl group, t-Bu represents a tertiary butyl group, c-Bu represents a  
15 cyclobutyl group, c-Pen represents a cyclopentyl group, c-Hex represents a cyclohexyl group, Ac represents an acetyl group, and Ph represents a phenyl group.

Table 1-1



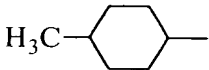
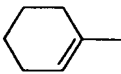
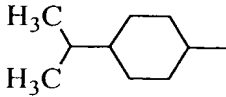
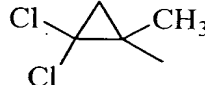
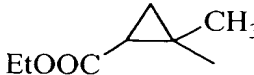
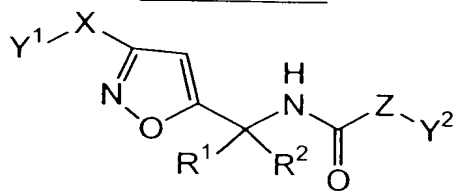
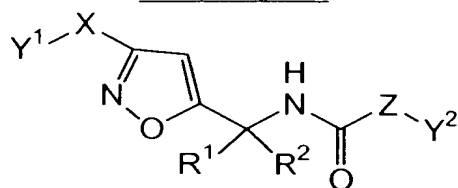
Compound No.	Y¹	X	R¹	R²	Z	Y²
I-1	Me-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-2	Et-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-3	n-Pr-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-4	i-Pr-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-5	c-Pr-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-6	c-Hex-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-7	n-Bu-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-8	i-Bu-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-9	t-Bu-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-10	Me <sub>2</sub> C=CH-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-11	I-C≡C-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-12	c-Pen-	—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-13		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-14		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-15		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-16		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-17		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -

Table 1-2



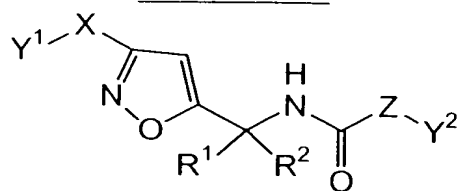
Compound No.	Y¹	X	R¹	R²	Z	Y²
I-18		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-19		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-20		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-21		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-22		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-23		—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-24	4-Me-3-F-C <sub>6</sub> H <sub>3</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-25	4-Me-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-26	4-MeO-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-27	4-F-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-28	2-Cl-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-29	3-Cl-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-30	4-Cl-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-31	2-pyridyl—	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-32	2-thienyl—	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-33	2-furyl—	—	H	H	O	C <sub>6</sub> H <sub>5</sub> —
I-34	4-Me-C <sub>6</sub> H <sub>4</sub> —	—	H	H	O	2-MeO-C <sub>6</sub> H <sub>4</sub> —

Table 1-3.



Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Z	Y <sup>2</sup>
I-35	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	H	H	O	2-Me-C <sub>6</sub> H <sub>4</sub> -
I-36	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	H	H	O	2-HOOC-C <sub>6</sub> H <sub>4</sub> -
I-37	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	H	H	O	2-MeOOC-C <sub>6</sub> H <sub>4</sub> -
I-38	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-39	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -CH <sub>2</sub> -	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-40	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-41	C <sub>6</sub> H <sub>5</sub> -	—	H	H	-O-CH <sub>2</sub> -	C <sub>6</sub> H <sub>5</sub> -
I-42	C <sub>6</sub> H <sub>5</sub> -	—	H	H	-O-CH <sub>2</sub> -	HC≡C—
I-43	C <sub>6</sub> H <sub>5</sub> -	—	H	H	-O-CH <sub>2</sub> -	I-C≡C—
I-44	C <sub>6</sub> H <sub>5</sub> -	—	H	H	-O-CH <sub>2</sub> -	H <sub>2</sub> C=CH=CH <sub>2</sub>
I-45	C <sub>6</sub> H <sub>5</sub> -	—	H	H		C <sub>6</sub> H <sub>5</sub> -
I-46	C <sub>6</sub> H <sub>5</sub> -	—	H	H		C <sub>6</sub> H <sub>5</sub> -
I-47	i-Pr-	—	H	H		C <sub>6</sub> H <sub>5</sub> -
I-48	C <sub>6</sub> H <sub>5</sub> -	—	H	H	-CH <sub>2</sub> -	C <sub>6</sub> H <sub>5</sub> -
I-49	C <sub>6</sub> H <sub>5</sub> -	—	H	H		C <sub>6</sub> H <sub>5</sub> -
I-50	C <sub>6</sub> H <sub>5</sub> -	—	H	H		c-Pr-

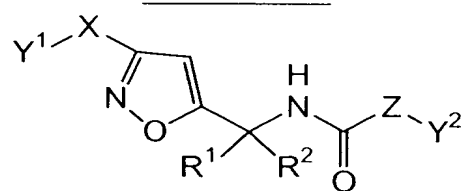
Table 1-4



Compound No.	Y¹	X	R¹	R²	Z	Y²
I-52	C <sub>6</sub> H <sub>5</sub> -		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-53		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-54		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-55		—	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-56	i-Pr-		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-57	t-Bu-		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-58	c-Pr-		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-59	i-Pr-		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-60	t-Bu-		H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-61	c-Pr-		H	H	O	C <sub>6</sub> H <sub>5</sub> -

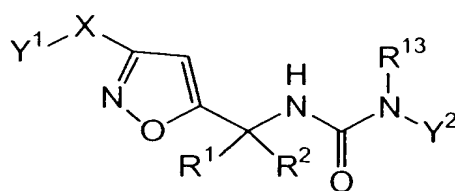


Table 1-5



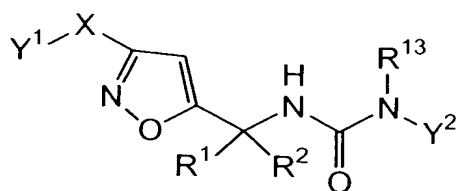
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Z	Y <sup>2</sup>
I-62	i-Pr-	-	H	H	-CH <sub>2</sub> -O-	C <sub>6</sub> H <sub>5</sub> -
I-63	i-Pr-	-	H	H	-CH <sub>2</sub> -	C <sub>6</sub> H <sub>5</sub> -
I-64	i-Bu-	-	H	H	-CH <sub>2</sub> -	C <sub>6</sub> H <sub>5</sub> -
I-65	ClCH <sub>2</sub> -	-	H	H	-CH <sub>2</sub> -O-	C <sub>6</sub> H <sub>5</sub> -
I-66	ClCH <sub>2</sub> -	-	H	H	O	C <sub>6</sub> H <sub>5</sub> -
I-67	ClCH <sub>2</sub> -	-	H	H	-CH <sub>2</sub> -	C <sub>6</sub> H <sub>5</sub> -

Table 2-1.



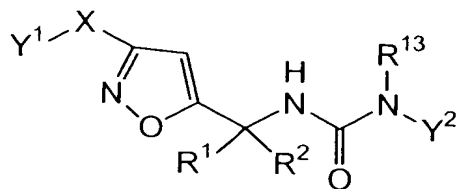
Compound No.	Y¹	X	R¹	R²	R¹³	Y²
II-1	4-Me-C₆H₄-	—	H	H	H	4-Me-C₆H₄-
II-2	4-Me-C₆H₄-	—	H	H	H	C₆H₅-
II-3	i-Pr-	—	H	H	H	C₆H₅-
II-4	i-Pr-	—	H	H	Me	C₆H₅-
II-5	C₆H₅-	—	H	H	H	c-Pr-
II-6	C₆H₅-	—	H	H	H	c-Pen-
II-7	C₆H₅-	—	H	H	H	c-Hex-
II-8	4-MeO-C₆H₄-	—	H	H	H	c-Hex-
II-9	4-Me-C₆H₄-	—	H	H	H	c-Hex-
II-10	4-Me-C₆H₄-	—	H	H	H	c-Pen-
II-11	C₆H₅-	—	H	H	H	
II-12	C₆H₅-	—	H	H	Me	C₆H₆-
II-13	C₆H₅-	-CH₂-	H	H	H	c-Hex-
II-14	C₆H₅-	-CH₂-	H	H	H	c-Pen-
II-15	i-Pr-	—	H	H	H	c-Hex-
II-16	C₆H₅-	—	H	H	H	
II-17	C₆H₅-	—	H	H	H	
II-18	4-Me-C₆H₄-	—	Me-	H	H	c-Hex-
II-19	4-Me-C₆H₄-	—	Et-	H	H	c-Hex-

Table 2-2.



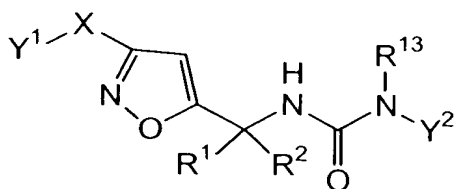
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	R <sup>13</sup>	Y <sup>2</sup>
II-20	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	c-Pr-	H	H	c-Hex-
II-21	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	I-Pr-	H	H	c-Hex-
II-22	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	MeO-	H	H	c-Hex-
II-23	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	MeOOC-	H	H	c-Hex-
II-24	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	NC-	H	H	c-Hex-
II-25	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Cl-	H	H	c-Hex-
II-26	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Me-	Me-	H	c-Hex-
II-27	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Me-	MeO-	H	c-Hex-
II-28	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Me-	MeOOC-	H	c-Hex-
II-29	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Me-	NC-	H	c-Hex-
II-30	4-Me-C <sub>6</sub> H <sub>4</sub> -	—	Me-	Cl-	H	c-Hex-
II-31	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-Me-C <sub>6</sub> H <sub>4</sub> -
II-32	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-Me-C <sub>6</sub> H <sub>4</sub> -
II-33	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
II-34	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-t-Bu-C <sub>6</sub> H <sub>4</sub> -
II-35	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-MeO-C <sub>6</sub> H <sub>4</sub> -
II-36	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-MeO-C <sub>6</sub> H <sub>4</sub> -
II-37	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
II-38	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-EtO-C <sub>6</sub> H <sub>4</sub> -
II-39	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-t-BuO-C <sub>6</sub> H <sub>4</sub> -
II-40	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-t-BuO-C <sub>6</sub> H <sub>4</sub> -
II-41	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-i-PrO-C <sub>6</sub> H <sub>4</sub> -

Table 2-3



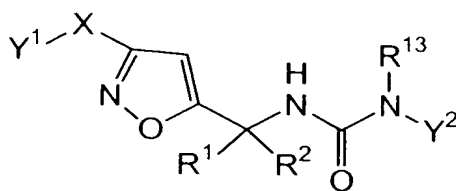
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	R <sup>13</sup>	Y <sup>2</sup>
II-42	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-PhO-C <sub>6</sub> H <sub>4</sub> -
II-43	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeS-C <sub>6</sub> H <sub>4</sub> -
II-44	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeO <sub>2</sub> S-C <sub>6</sub> H <sub>4</sub> -
II-45	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-F-C <sub>6</sub> H <sub>4</sub> -
II-46	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-F-C <sub>6</sub> H <sub>4</sub> -
II-47	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-F-C <sub>6</sub> H <sub>4</sub> -
II-48	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-Cl-C <sub>6</sub> H <sub>4</sub> -
II-49	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-Cl-C <sub>6</sub> H <sub>4</sub> -
II-50	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
II-51	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-Br-C <sub>6</sub> H <sub>4</sub> -
II-52	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-Br-C <sub>6</sub> H <sub>4</sub> -
II-53	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Br-C <sub>6</sub> H <sub>4</sub> -
II-54	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
II-55	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
II-56	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
II-57	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-NC-C <sub>6</sub> H <sub>4</sub> -
II-58	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-NC-C <sub>6</sub> H <sub>4</sub> -
II-59	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-NC-C <sub>6</sub> H <sub>4</sub> -
II-60	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2-HO-C <sub>6</sub> H <sub>4</sub> -
II-61	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-HO-C <sub>6</sub> H <sub>4</sub> -
II-62	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-HO-C <sub>6</sub> H <sub>4</sub> -
II-63	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -

Table 2-4



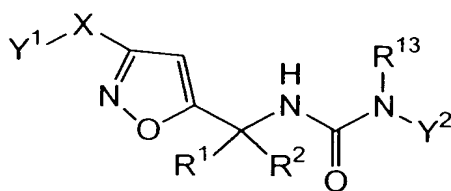
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	R <sup>13</sup>	Y <sup>2</sup>
II-64	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-CF <sub>3</sub> S-C <sub>6</sub> H <sub>4</sub> -
II-65	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-CF <sub>3</sub> O-C <sub>6</sub> H <sub>4</sub> -
II-66	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-F <sub>2</sub> HC-O-C <sub>6</sub> H <sub>4</sub> -
II-67	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-HOOC-C <sub>6</sub> H <sub>4</sub> -
II-68	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -
II-69	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-MeHNOC-C <sub>6</sub> H <sub>4</sub> -
II-70	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeHNOC-C <sub>6</sub> H <sub>4</sub> -
II-71	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-MeOCHN-C <sub>6</sub> H <sub>4</sub> -
II-72	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeOCHN-C <sub>6</sub> H <sub>4</sub> -
II-73	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
II-74	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
II-75	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
II-76	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
II-77	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Ph-C <sub>6</sub> H <sub>4</sub> -
II-78	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2,5-(Cl) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
II-79	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3,4,(F) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
II-80	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3,4,-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
II-81	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	2,5-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
II-82	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3,4-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
II-83	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	3-Cl-4-MeO-C <sub>6</sub> H <sub>3</sub> -
II-84	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-MeO-2-HO-C <sub>6</sub> H <sub>3</sub> -
II-85	C <sub>6</sub> H <sub>5</sub> -	—	H	H	H	4-Me-2-HO-C <sub>6</sub> H <sub>3</sub> -

Table 2-5



Compound No.	Y¹	X	R¹	R²	R¹³	Y²
II-86	C₆H₅-	—	H	H	H	4-MeO-3-F-C₆H₃-
II-87	C₆H₅-	—	H	H	H	3-F-4-MeO-C₆H₃-
II-88	C₆H₅-	—	H	H	H	4-Me-3-F-C₆H₃-
II-89	C₆H₅-	—	H	H	H	4-Me-3-Cl-C₆H₃-
II-90	C₆H₅-	—	H	H	H	4-Me-3-CF₃-C₆H₃-
II-91	C₆H₅-	—	H	H	H	4-ClCH₂-C₆H₃-
II-92	C₆H₅-	—	H	H	H	4-OHC-C₆H₄-
II-93	C₆H₅-	—	H	H	H	4-MeOOCHNHN-C₆H₄-
II-94	C₆H₅-	—	H	H	H	4-CN-3-F-C₆H₃-
II-95	C₆H₅-	—	H	H	H	4-Me-C₆H₄-
II-96	C₆H₅-	—	H	H	H	4-MeO-C₆H₄-
II-97	C₆H₅-	—	H	H	H	4-CF₃O-C₆H₄-
II-98	C₆H₅-	—	H	H	H	4-t-Bu-C₆H₄-
II-99	C₆H₅-	—	H	H	H	2-pyridyl-
II-100	C₆H₅-	—	H	H	H	3-pyridyl-
II-101	C₆H₅-	—	H	H	H	4-pyridyl-
II-102	C₆H₅-	—	H	H	H	2-thienyl-
II-103	C₆H₅-	—	H	H	H	2-furyl-

Table 2-6



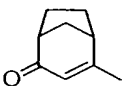
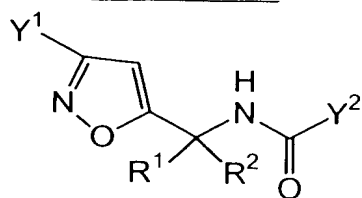
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	R <sup>13</sup>	Y <sup>2</sup>
II-104	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	H	c-Hex-
II-105	ClCH <sub>2</sub> -	-	H	H	H	c-Hex-
II-106	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	H	c-Pen-
II-107	ClCH <sub>2</sub> -	-	H	H	H	c-Pen-
II-108	Me-	-	H	H	H	c-Hex-
II-109	ClCH <sub>2</sub> -	-	H	H	H	C <sub>6</sub> H <sub>5</sub> -
II-110	Me-	-NHCO-	H	H	H	c-Hex-
II-111	BrCH <sub>2</sub> -	-	H	H	H	c-Hex-
II-112		-O-CH <sub>2</sub> -	H	H	H	c-Hex-
II-113	NC-CH <sub>2</sub> -	-	H	H	H	c-Hex-
II-114	MeS-CH <sub>2</sub> -	-	H	H	H	c-Hex-
II-115	ICH <sub>2</sub> -	-	H	H	H	c-Hex-
II-116	ClCH <sub>2</sub> -	-	H	H	Me	C <sub>6</sub> H <sub>5</sub> -
II-117	MeOCH <sub>2</sub> -	-	H	H	H	c-Hex-
II-118	CH <sub>3</sub> COOCH <sub>2</sub> -	-	H	H	H	c-Hex-
II-119	HOCH <sub>2</sub> -	-	H	H	H	c-Hex-

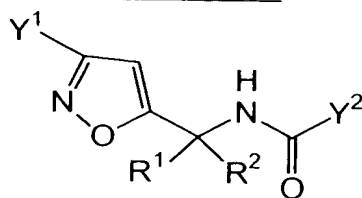
Table 3-1.



Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-1	Me-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-2	ClCH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-3	ClCH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-4	i-Pr-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-5	i-Pr-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-6	c-Hex-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-7	2-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	2-HO-C <sub>6</sub> H <sub>4</sub> -
III-8	2-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-9	3-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-10	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-11	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-12	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-13	4-Et-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-14	4-t-Bu-C <sub>6</sub> H <sub>4</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-15	2-MeO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-16	3-MeO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-17	4-MeO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-18	4-MeO-C <sub>6</sub> H <sub>4</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-19	4-MeO-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-20	4-EtO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-21	4-t-BuO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-22	3-t-BuO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-23	4-i-PrO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-24	4-PhO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -

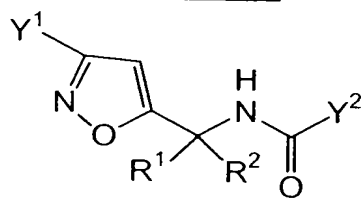


Table 3-2.



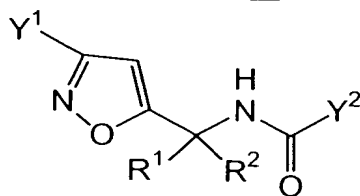
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-25	4-MeS-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-26	4-MeO <sub>2</sub> S-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-27	4-MeOS-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-28	4-MeS-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-29	2-F-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-30	3-F-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-31	4-F-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-32	2-Cl-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-33	3-Cl-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-34	4-Cl-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-35	2-Br-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-36	3-Br-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-37	4-Br-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-38	2-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-39	3-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-40	4-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-41	2-NC-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-42	3-NC-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-43	4-NC-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-44	2-HO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-45	3-HO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-46	4-HO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-47	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-48	4-CF <sub>3</sub> S-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -

Table 3-3.



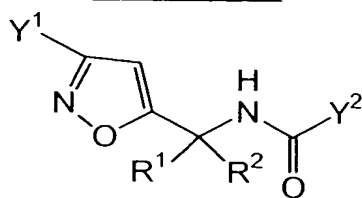
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-49	4-CF <sub>3</sub> O-C <sub>6</sub> H <sub>4</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-50	4-H <sub>2</sub> FCO-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-51	4-HOOC-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-52	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-53	3-MeHNOC-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-54	4-MeHNOC-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-55	4-MeOHNOC-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-56	3-MeOCHN-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-57	4-MeOCHN-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-58	3-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-59	4-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-60	3-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-61	4-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-62	4-Ph-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-63	2,5-(Cl) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-64	3,4,(F) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-65	2,4,-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-66	2,4-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-67	3,4-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-68	3,4-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-69	3-Cl-4-MeO-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-70	4-MeO-2-HO-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-71	4-Me-2-HO-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-72	4-MeO-3-F-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -

Table 3-4.



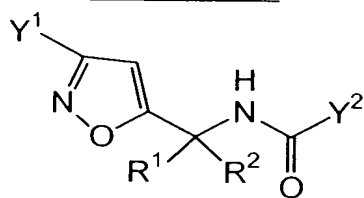
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-73	4-Me-3-F-C <sub>6</sub> H <sub>3</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-74	4-MeO-2-F-C <sub>6</sub> H <sub>3</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-75	3-F-4-MeO-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-76	4-Me-3-F-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-77	4-Me-3-Cl-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-78	4-Me-3-CF <sub>3</sub> -C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-79	4-ClCH <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-80	4-OHC-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-81	4-MeOOCHNHN-C <sub>6</sub> H <sub>4</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-82	4-CN-3-F-C <sub>6</sub> H <sub>3</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-83	2-pyridyl-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-84	3-pyridyl-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-85	4-pyridyl-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-86	2-thienyl-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-87	2-furyl-	H	H	C <sub>6</sub> H <sub>5</sub> -
III-88		H	H	C <sub>6</sub> H <sub>5</sub> -
III-89		H	H	C <sub>6</sub> H <sub>5</sub> -
III-90		H	H	C <sub>6</sub> H <sub>5</sub> -
III-91		H	H	C <sub>6</sub> H <sub>5</sub> -

Table 3-5



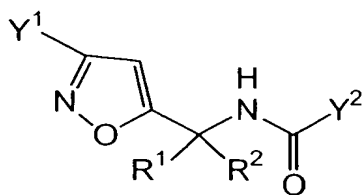
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-92		H	H	C <sub>6</sub> H <sub>5</sub> -
III-93		H	H	C <sub>6</sub> H <sub>5</sub> -
III-94		H	H	C <sub>6</sub> H <sub>5</sub> -
III-95		H	H	C <sub>6</sub> H <sub>5</sub> -
III-96		H	H	C <sub>6</sub> H <sub>5</sub> -
III-97		H	H	C <sub>6</sub> H <sub>5</sub> -
III-98		H	H	C <sub>6</sub> H <sub>5</sub> -
III-99		H	H	C <sub>6</sub> H <sub>5</sub> -

Table 3-6



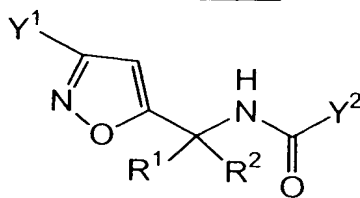
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-100		H	H	C <sub>6</sub> H <sub>5</sub> -
III-101		H	H	C <sub>6</sub> H <sub>5</sub> -
III-102		H	H	C <sub>6</sub> H <sub>5</sub> -
III-103		H	H	C <sub>6</sub> H <sub>5</sub> -
III-104		H	H	C <sub>6</sub> H <sub>5</sub> -
III-105	C <sub>6</sub> H <sub>5</sub> -	H	H	2-Me-C <sub>6</sub> H <sub>4</sub> -
III-106	C <sub>6</sub> H <sub>5</sub> -	H	H	3-Me-C <sub>6</sub> H <sub>4</sub> -
III-107	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-108	C <sub>6</sub> H <sub>5</sub> -	H	H	4-t-Bu-C <sub>6</sub> H <sub>4</sub> -
III-109	C <sub>6</sub> H <sub>5</sub> -	H	H	2-MeO-C <sub>6</sub> H <sub>4</sub> -
III-110	C <sub>6</sub> H <sub>5</sub> -	H	H	3-MeO-C <sub>6</sub> H <sub>4</sub> -
III-111	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-112	C <sub>6</sub> H <sub>5</sub> -	H	H	4-EtO-C <sub>6</sub> H <sub>4</sub> -

Table 3-7.



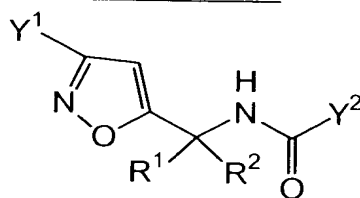
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-113	C <sub>6</sub> H <sub>5</sub> -	H	H	4-t-BuO-C <sub>6</sub> H <sub>4</sub> -
III-114	C <sub>6</sub> H <sub>5</sub> -	H	H	3-t-BuO-C <sub>6</sub> H <sub>4</sub> -
III-115	C <sub>6</sub> H <sub>5</sub> -	H	H	4-i-PrO-C <sub>6</sub> H <sub>4</sub> -
III-116	C <sub>6</sub> H <sub>5</sub> -	H	H	4-PhO-C <sub>6</sub> H <sub>4</sub> -
III-117	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeS-C <sub>6</sub> H <sub>4</sub> -
III-118	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeO <sub>2</sub> S-C <sub>6</sub> H <sub>4</sub> -
III-119	C <sub>6</sub> H <sub>5</sub> -	H	H	2-F-C <sub>6</sub> H <sub>4</sub> -
III-120	C <sub>6</sub> H <sub>5</sub> -	H	H	3-F-C <sub>6</sub> H <sub>4</sub> -
III-121	C <sub>6</sub> H <sub>5</sub> -	H	H	4-F-C <sub>6</sub> H <sub>4</sub> -
III-122	C <sub>6</sub> H <sub>5</sub> -	H	H	2-Cl-C <sub>6</sub> H <sub>4</sub> -
III-123	C <sub>6</sub> H <sub>5</sub> -	H	H	3-Cl-C <sub>6</sub> H <sub>4</sub> -
III-124	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-125	C <sub>6</sub> H <sub>5</sub> -	H	H	2-Br-C <sub>6</sub> H <sub>4</sub> -
III-126	C <sub>6</sub> H <sub>5</sub> -	H	H	3-Br-C <sub>6</sub> H <sub>4</sub> -
III-127	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Br-C <sub>6</sub> H <sub>4</sub> -
III-128	C <sub>6</sub> H <sub>5</sub> -	H	H	2-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
III-129	C <sub>6</sub> H <sub>5</sub> -	H	H	3-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
III-130	C <sub>6</sub> H <sub>5</sub> -	H	H	4-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
III-131	C <sub>6</sub> H <sub>5</sub> -	H	H	2-NC-C <sub>6</sub> H <sub>4</sub> -
III-132	C <sub>6</sub> H <sub>5</sub> -	H	H	3-NC-C <sub>6</sub> H <sub>4</sub> -
III-133	C <sub>6</sub> H <sub>5</sub> -	H	H	4-NC-C <sub>6</sub> H <sub>4</sub> -
III-134	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	2-HO-C <sub>6</sub> H <sub>4</sub> -
III-135	C <sub>6</sub> H <sub>5</sub> -	H	H	3-HO-C <sub>6</sub> H <sub>4</sub> -
III-136	C <sub>6</sub> H <sub>5</sub> -	H	H	4-HO-C <sub>6</sub> H <sub>4</sub> -

Table 3-8.



Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-137	C <sub>6</sub> H <sub>5</sub> -	H	H	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -
III-138	C <sub>6</sub> H <sub>5</sub> -	H	H	4-CF <sub>3</sub> S-C <sub>6</sub> H <sub>4</sub> -
III-139	C <sub>6</sub> H <sub>5</sub> -	H	H	4-CF <sub>3</sub> O-C <sub>6</sub> H <sub>4</sub> -
III-140	C <sub>6</sub> H <sub>5</sub> -	H	H	4-F <sub>2</sub> HC-O-C <sub>6</sub> H <sub>4</sub> -
III-141	C <sub>6</sub> H <sub>5</sub> -	H	H	4-HOOC-C <sub>6</sub> H <sub>4</sub> -
III-142	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -
III-143	C <sub>6</sub> H <sub>5</sub> -	H	H	3-MeHNOC-C <sub>6</sub> H <sub>4</sub> -
III-144	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeHNOC-C <sub>6</sub> H <sub>4</sub> -
III-145	C <sub>6</sub> H <sub>5</sub> -	H	H	3-MeOCHN-C <sub>6</sub> H <sub>4</sub> -
III-146	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeOCHN-C <sub>6</sub> H <sub>4</sub> -
III-147	C <sub>6</sub> H <sub>5</sub> -	H	H	3-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
III-148	C <sub>6</sub> H <sub>5</sub> -	H	H	4-H <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
III-149	C <sub>6</sub> H <sub>5</sub> -	H	H	3-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
III-150	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Me <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -
III-151	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Ph-C <sub>6</sub> H <sub>4</sub> -
III-152	C <sub>6</sub> H <sub>5</sub> -	H	H	2,5-(Cl) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-153	C <sub>6</sub> H <sub>5</sub> -	H	H	3,4,(F) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-154	C <sub>6</sub> H <sub>5</sub> -	H	H	3,4,-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-155	C <sub>6</sub> H <sub>5</sub> -	H	H	2,5-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-156	C <sub>6</sub> H <sub>5</sub> -	H	H	3,4-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-157	C <sub>6</sub> H <sub>5</sub> -	H	H	3-Cl-4-MeO-C <sub>6</sub> H <sub>3</sub> -
III-158	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeO-2-HO-C <sub>6</sub> H <sub>3</sub> -
III-159	C <sub>6</sub> H <sub>5</sub> -	H	H	4-Me-2-HO-C <sub>6</sub> H <sub>3</sub> -
III-160	C <sub>6</sub> H <sub>5</sub> -	H	H	4-MeO-3-F-C <sub>6</sub> H <sub>3</sub> -

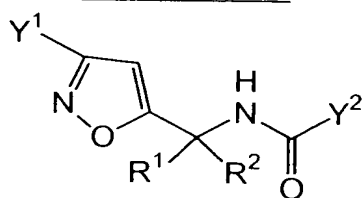
Table 3-9



Compound No.	Y¹	R¹	R²	Y²
III-161	C₆H₅-	H	H	3-F-4-MeO-C₆H₃-
III-162	4-Me-C₆H₄-	H	H	4-Me-3-F-C₆H₃-
III-163	C₆H₅-	H	H	4-Me-3-Cl-C₆H₃-
III-164	C₆H₅-	H	H	4-Me-3-CF₃-C₆H₃-
III-165	C₆H₅-	H	H	4-ClCH₂-C₆H₄-
III-166	C₆H₅-	H	H	4-OHC-C₆H₄-
III-167	C₆H₅-	H	H	4-MeOOCHNHN-C₆H₄-
III-168	C₆H₅-	H	H	2-HO-3-OHCHN-C₆H₃-
III-169	C₆H₅-	H	H	4-CN-3-F-C₆H₃-
III-170	C₆H₅-	H	H	4-Me-C₆H₄-
III-171	C₆H₅-	H	H	4-MeO-C₆H₄-
III-172	C₆H₅-	H	H	4-CF₃O-C₆H₄-
III-173	C₆H₅-	H	H	4-t-Bu-C₆H₄-
III-174	C₆H₅-	H	H	2-pyridyl-
III-175	C₆H₅-	H	H	3-pyridyl-
III-176	C₆H₅-	H	H	4-pyridyl-
III-177	C₆H₅-	H	H	2-thienyl-
III-178	C₆H₅-	H	H	2-furyl-
III-180	C₆H₅-	H	H	
III-181	C₆H₅-	H	H	

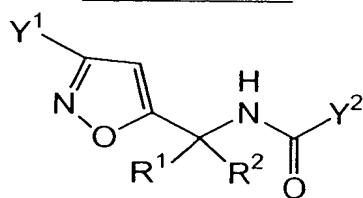


Table 3-10



Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-182	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-183	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-184	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-185	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-186	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-187	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-188	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-189	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-190	C <sub>6</sub> H <sub>5</sub> -	H	H	

Table 3-11.



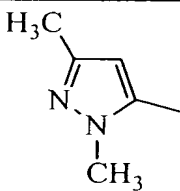
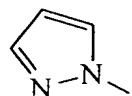
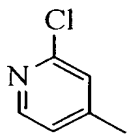
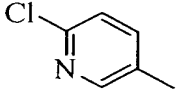
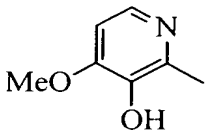
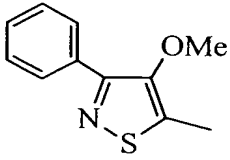
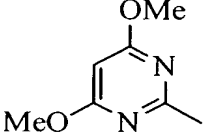
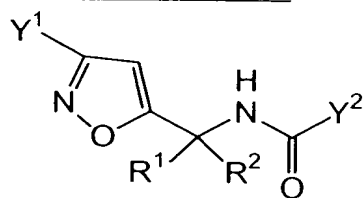
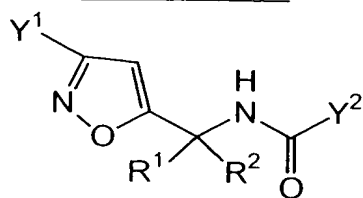
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-191	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-192	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-193	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-194	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-195	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-196	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-197	C <sub>6</sub> H <sub>5</sub> -	H	H	

Table 3-12



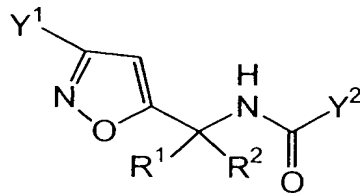
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-198	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-199	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-200	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-201	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-202	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-203	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-204	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-205	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-206	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	

Table 3-13.



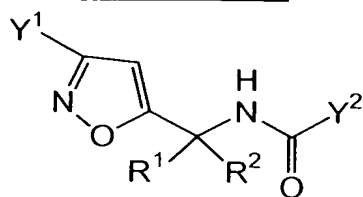
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-207	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-208	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-209	C <sub>6</sub> H <sub>5</sub> -	H	H	Me <sub>2</sub> C=CH-
III-210		H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-211		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -

Table 3-14



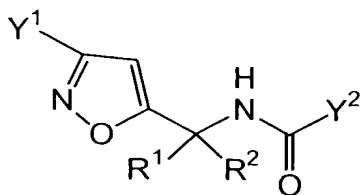
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-212	C <sub>6</sub> H <sub>5</sub> -	H	H	
III-213	C <sub>6</sub> H <sub>5</sub> -	Me	H	
III-214		H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-215	4-MeO-2-F-C <sub>6</sub> H <sub>3</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-216	Me <sub>2</sub> C=CH-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-217	2-pyridyl-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-218	3-pyridyl-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-219	4-pyridyl-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-220	3-thienyl-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-221		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-222	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	c-Hex-
III-223		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-224		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-225	i-Pr-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-226	ClCH <sub>2</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
III-227	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	c-Pen-
III-228	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	c-Bu-

Table 3-15



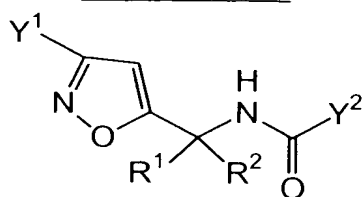
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-229	i-Bu-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-230		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-231	2-Cl-C <sub>6</sub> H <sub>4</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-232		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-233	ClCH <sub>2</sub> -	H	H	c-Hex-
III-234	i-Pr-	H	H	c-Hex-
III-235	n-Bu-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-236		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-237	t-Bu-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-238	c-Hex-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-239		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-240	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-241	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	
III-242	4-Me-C <sub>6</sub> H <sub>4</sub> -	H	H	

Table 3-16



Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-243	i-Pr-	H	H	
III-244	ClCH <sub>2</sub> -	H	H	
III-245	n-Pr-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-246		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-247		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-248		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-249		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-250		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
III-251	i-Bu-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
III-252	i-Bu-	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-253	i-Bu-	H	H	2,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-254	ClCH <sub>2</sub> -	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-255	i-Bu-	H	H	3,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-256	i-Bu-	H	H	4-Br-C <sub>6</sub> H <sub>4</sub> -
III-257	i-Bu-	H	H	4-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -

Table 3-17



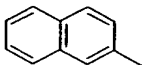
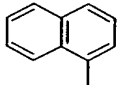
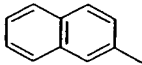
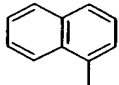
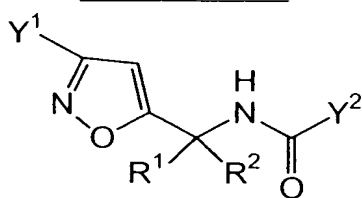
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-258	i-Bu-	H	H	4-Et-C <sub>6</sub> H <sub>4</sub> -
III-259	i-Bu-	H	H	
III-260	ClCH <sub>2</sub> -	H	H	4-Br-C <sub>6</sub> H <sub>4</sub> -
III-261	ClCH <sub>2</sub> -	H	H	4-CF <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -
III-262	ClCH <sub>2</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -CH=CH-
III-263	i-Bu-	H	H	
III-264	n-Pr-	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-265	n-Pr-	H	H	4-Br-C <sub>6</sub> H <sub>4</sub> -
III-266	n-Pr-	H	H	2,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-267	ClCH <sub>2</sub> -	H	H	
III-268	ClCH <sub>2</sub> -	H	H	
III-269	ClCH <sub>2</sub> -	H	H	4-Et-C <sub>6</sub> H <sub>4</sub> -
III-270	ClCH <sub>2</sub> -	H	H	4-ClCH <sub>2</sub> O-C <sub>6</sub> H <sub>4</sub> -
III-271	ClCH <sub>2</sub> -	H	H	2,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-272	ClCH <sub>2</sub> -	H	H	3,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-273	ClCH <sub>2</sub> -	H	H	2-Me-C <sub>6</sub> H <sub>4</sub> -
III-274	ClCH <sub>2</sub> -	H	H	3-Me-C <sub>6</sub> H <sub>4</sub> -
III-275	BrCH <sub>2</sub> -	H	H	2,4-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-276	ClCH <sub>2</sub> -	H	H	2-Cl-C <sub>6</sub> H <sub>4</sub> -
III-277	ClCH <sub>2</sub> -	H	H	3-Cl-C <sub>6</sub> H <sub>4</sub> -

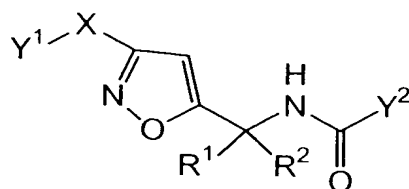


Table 3-18 .



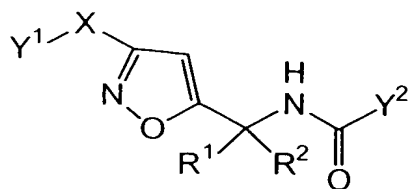
Compound No.	Y <sup>1</sup>	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
III-278	ClCH <sub>2</sub> -	H	H	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -
III-279	ClCH <sub>2</sub> -	H	H	2-Cl-4-SO <sub>2</sub> Me-C <sub>6</sub> H <sub>3</sub> -
III-280	ClCH <sub>2</sub> -	H	H	2-thienyl-
III-281	ClCH <sub>2</sub> -	H	H	4-F-C <sub>6</sub> H <sub>4</sub> -
III-282	ClCH <sub>2</sub> -	H	H	4-Ph-C <sub>6</sub> H <sub>4</sub> -
III-283	ClCH <sub>2</sub> -	H	H	4-NC-C <sub>6</sub> H <sub>4</sub> -
III-284	HOCH <sub>2</sub> -	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-285		H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-286	ClCH <sub>2</sub> -	H	H	2-F-C <sub>6</sub> H <sub>4</sub> -
III-287	ClCH <sub>2</sub> -	H	H	3-F-C <sub>6</sub> H <sub>4</sub> -
III-288	ClCH <sub>2</sub> -	H	H	4-I-C <sub>6</sub> H <sub>4</sub> -
III-289	ClCH <sub>2</sub> -	H	H	4-n-Pr-C <sub>6</sub> H <sub>4</sub> -
III-290	ClCH <sub>2</sub> -	H	H	4-SMe-C <sub>6</sub> H <sub>4</sub> -
III-291	ClCH <sub>2</sub> -	H	H	4-SO <sub>2</sub> Me-C <sub>6</sub> H <sub>4</sub> -
III-292	ClCH <sub>2</sub> -	H	H	2,6-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-293	ClCH <sub>2</sub> -	H	H	3,5-Cl <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -
III-294	ClCH <sub>2</sub> -	H	H	2,3,4-F <sub>3</sub> -C <sub>6</sub> H <sub>2</sub> -
III-295	ClCH <sub>2</sub> -	H	H	2,4,5-F <sub>3</sub> -C <sub>6</sub> H <sub>2</sub> -
III-296	ClCH <sub>2</sub> -	H	H	2-pyridyl-
III-297	ClCH <sub>2</sub> -	H	H	4-pyridyl-
III-298	FCH <sub>2</sub> -	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
III-299	ClCH <sub>2</sub> -	H	H	2,3,6-F <sub>3</sub> -C <sub>6</sub> H <sub>2</sub> -

Table 4-1'



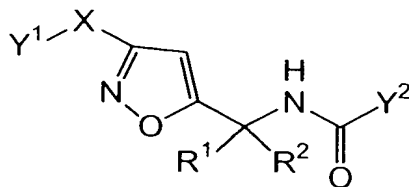
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-1	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
IV-2	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-3	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-4	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	c-Hex-
IV-5	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	c-Hexyloxy-
IV-6	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	4-Et-C <sub>6</sub> H <sub>4</sub> -
IV-7	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	2-HO-C <sub>6</sub> H <sub>4</sub> -
IV-8	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	4-MeO <sub>2</sub> S-C <sub>6</sub> H <sub>4</sub> -
IV-9	4-MeO-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-10	4-Me-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-11	3-Me-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-12	2-Me-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-13	4-Me-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-14	4-NC-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-15	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-16	3-F-4-MeO-C <sub>6</sub> H <sub>3</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-17	4-Me-3-F-C <sub>6</sub> H <sub>3</sub> -	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-18	2-pyridyl-	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-19	3-pyridyl-	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-20	4-pyridyl-	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-21	2-thienyl-	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-22	2-furyl-	-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-23	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -	H	H	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -

Table 4-2.



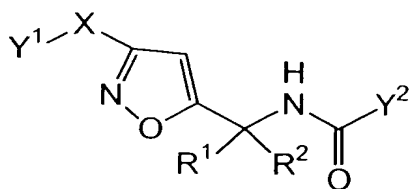
Compound No.	Y¹	X	R¹	R²	Y²
IV-24		-CH₂-	H	H	4-MeO-C₆H₄-
IV-25		-CH₂-	H	H	4-MeO-C₆H₄-
IV-26		-CH₂-	H	H	4-MeO-C₆H₄-
IV-27		-CH₂-	H	H	4-MeO-C₆H₄-
IV-28		-CH₂-	H	H	4-MeO-C₆H₄-
IV-29	C₆H₅-		H	H	C₆H₅-
IV-30	C₆H₅-		H	H	C₆H₅-
IV-31	C₆H₅-		H	H	C₆H₅-
IV-32	C₆H₅-		H	H	C₆H₅-
IV-33	C₆H₅-	-CH₂-CH₂-	H	H	C₆H₅-
IV-34	C₆H₅-	-CH₂-CH₂-	H	H	4-MeO-C₆H₄-
IV-35	C₆H₅-	-CH₂-CH₂-	H	H	4-Me-C₆H₄-

Table 4-3



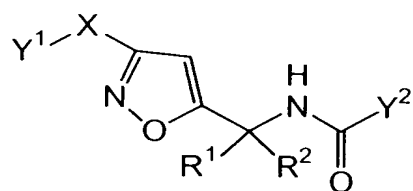
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-36	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-37	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-38	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-39	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-40	C <sub>6</sub> H <sub>5</sub> -		H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-41	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-42	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-43	C <sub>6</sub> H <sub>5</sub> -		H	H	C <sub>6</sub> H <sub>5</sub> -
IV-44	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	C <sub>6</sub> H <sub>5</sub> -
IV-45	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-46	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -

Table 4-4.



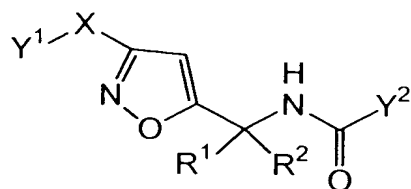
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-47	2-Me-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-48	3-Me-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-49	3-Me-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-50	4-Me-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-51	2-NC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-52	3-NC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-53	4-NC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-54	2-MeO-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-55	3-MeO-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-56	4-MeO-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-57	2-F-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-58	3-F-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-59	4-F-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-60	4-O <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-61	c-Hex-	-O-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-62	2-pyridyl-	-O-CH <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-63	C <sub>6</sub> H <sub>5</sub> -	-O-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-64	C <sub>6</sub> H <sub>5</sub> -	-S-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-65	C <sub>6</sub> H <sub>5</sub> -	-SO <sub>2</sub> -	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-66	C <sub>6</sub> H <sub>5</sub> -	-NH-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-67	C <sub>6</sub> H <sub>5</sub> -		H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-68	C <sub>6</sub> H <sub>5</sub> -	-CH <sub>2</sub> -O-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -

Table 4-5



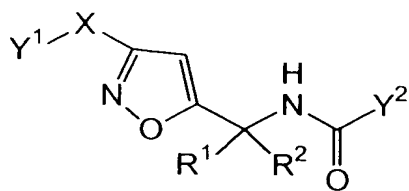
Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-69	C <sub>6</sub> H <sub>5</sub> -		H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-70	C <sub>6</sub> H <sub>5</sub> -	-CO-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-71	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-72		-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-73		-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-74		-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -

Table 4-6



Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-75		-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-76	2,6-(MeO) <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-77		-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-78		-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-79		-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-80	2,6-(Me) <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-81	4-MeOCHN-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-82	2-MeOOC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-83	3-MeOOC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-84	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-85	4-MeOOC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-86	2,6-F <sub>2</sub> -C <sub>6</sub> H <sub>3</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-87	4-Ac-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-88	4-MeHNOC-C <sub>6</sub> H <sub>4</sub> -	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-89	2-furyl-		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-90	4-pyridyl-	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -

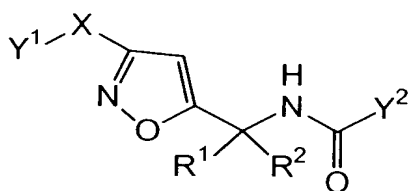
Table 4-7



Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-91	3-pyridyl-	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-92	2-pyridyl-	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-93	4-F <sub>2</sub> HCO-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-94	4-MeO-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-95	4-F-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-96	2-Cl-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-97	4-NC-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-98	C <sub>6</sub> H <sub>5</sub> -	-CO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-99	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	
IV-100	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-101	4-MeO-C <sub>6</sub> H <sub>4</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-102	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-103	Me-		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-104	i-Pr-		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -



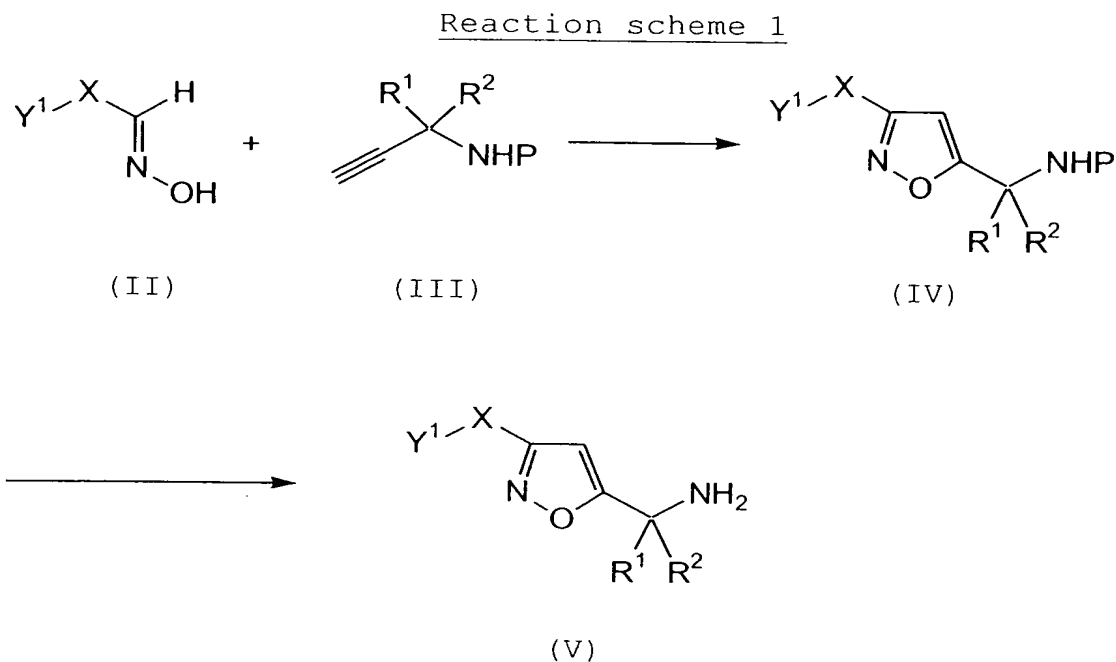
Table 4-8.



Compound No.	Y <sup>1</sup>	X	R <sup>1</sup>	R <sup>2</sup>	Y <sup>2</sup>
IV-105	3-Cl-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-106	4-Cl-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-107	2,4-F <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-108	4-Me-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-109	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-110	C <sub>6</sub> H <sub>5</sub> -	-C≡C-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-111	Me-	-O-CH <sub>2</sub> -	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-112	2-NC-C <sub>6</sub> H <sub>4</sub> -	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-113	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	C <sub>6</sub> H <sub>5</sub> -
IV-114	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	4-Cl-C <sub>6</sub> H <sub>4</sub> -
IV-115	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	4-Me-C <sub>6</sub> H <sub>4</sub> -
IV-116	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	3-Me-C <sub>6</sub> H <sub>4</sub> -
IV-117	C <sub>6</sub> H <sub>5</sub> -	-NHCO-	H	H	2-Me-C <sub>6</sub> H <sub>4</sub> -
IV-118	i-Pr-	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-119	t-Bu-	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-120	c-Pen-	-NHCO-	H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-121	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	c-Hex-
IV-122	C <sub>6</sub> H <sub>5</sub> -	-O-CH <sub>2</sub> -	H	H	
IV-123	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -
IV-124	C <sub>6</sub> H <sub>5</sub> -		H	H	4-MeO-C <sub>6</sub> H <sub>4</sub> -

[Method of manufacturing the compound of the invention]

[1] A substituted 5-alkylamino isoxazole derivative (V) which is an intermediate of the compound of the invention can be prepared by a method shown in the following  
5 reaction scheme (1).



(In the reaction scheme, P represents a protecting group of an amino group, X,  $Y^1$ ,  $R^1$ , and  $R^2$  represent the same meanings  
10 as those defined above.)

That is, a protection body of a substituted 5-alkylamino isoxazole derivative represented by the formula (IV) can be prepared by reacting an oxime derivative represented by the formula (II) with a protected propargyl  
15 amine derivative represented by the formula (III) in an appropriate solvent in the presence of Hypolites.

Here, a t-butoxycarbonyl group, a benzyloxycarbonyl

group, etc. are used as a protective group of an amino group.

Here, examples of Hypolites include: alkali metals such as sodium hypochlorite, potassium hypochlorite, lithium hypochlorite, sodium hypobromite, and potassium hypobromite; 5 calcium hypochlorite; magnesium hypochlorite; strontium hypochlorite; calcium hypobromite; and magnesium hypobromite. Preferable examples thereof include alkali metal solutions of sodium hypochlorite and potassium hypochlorite.

Preferable examples of the solvent used for the 10 reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, 15 carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; esters such as methyl acetate, ethyl acetate, and propyl acetate; water; acetonitrile; dimethyl formamide; dimethyl sulfoxide; N-methyl-2-pyrrolidone; and arbitrary mixed solvents thereof. More preferable examples 20 thereof include: halogenated hydrocarbons such as chloroform and dichloromethane; and arbitrary mixed solvents selected therefrom.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

25 Although the reaction time depends on the reaction

temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 3 days.

The compound of the formula (IV) which is a reaction intermediate is collected from a reaction mixture by a usual  
5 post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; washing with water; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

10 Subsequently, a substituted 5-aminoalkyl isoxazole derivative represented by the formula (V) can be prepared by reacting the compound of the formula (IV) in an appropriate solvent under an appropriate deprotection condition.

The deprotection condition used here depends on the  
15 protecting group. For example, when a t-butoxycarbonyl group is used as the protecting group, proton acid solutions such as an aqueous solution of hydrochloric acid are preferable, and when a benzyloxycarbonyl group is used as the protecting group, it is preferable to perform  
20 deprotection under a hydrogen flow in the presence of a catalyst such as palladium carbon or Raney Nickel.

Preferable examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene,  
25 and xylene; primary alcohols such as methanol, ethanol, and

n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; esters such as methyl acetate, ethyl acetate, and propyl acetate; water; acetonitrile; dimethyl formamide; dimethyl sulfoxide; and N-methyl-2-pyrrolidone; and arbitrary mixed solvents thereof.

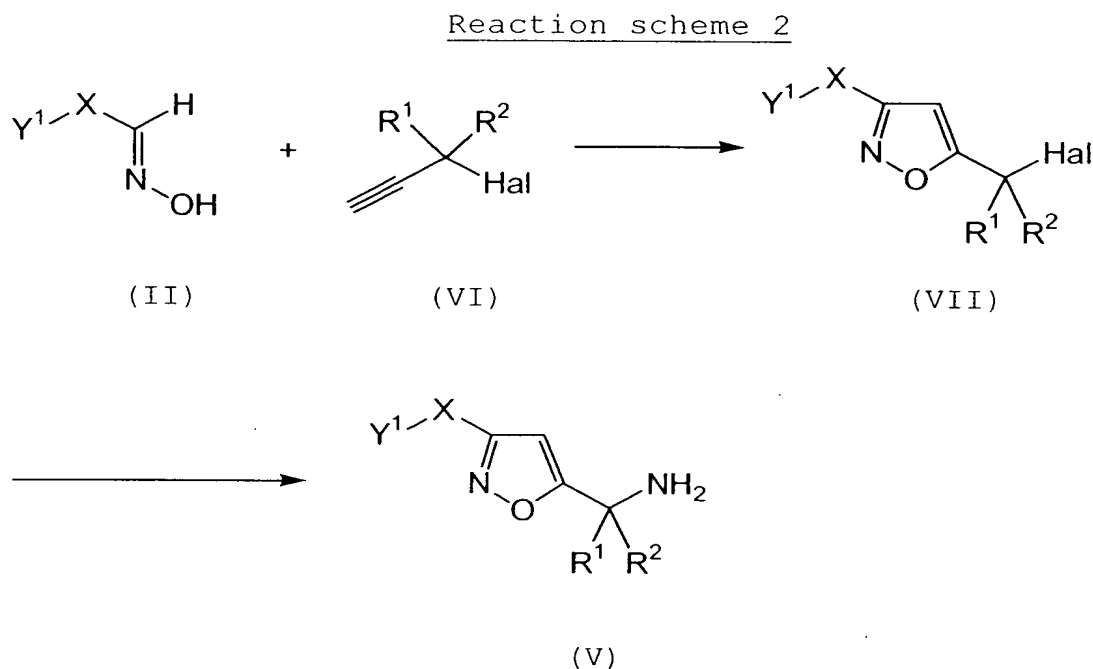
The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature, the reaction substrate, and the reaction reagent, the reaction is generally completed in 30 minutes to 1 week.

The aminoalkyl isoxazole derivative of the formula (V) which is the reaction intermediate is collected from the reaction mixture by the usual post-treatment. For example, in the case of the deprotection condition of the t-butoxycarbonyl group, the derivative is obtained by: making the reaction mixture alkali; then extracting with a solvent; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary. Also in the case of the deprotection condition of the benzyloxycarbonyl group, the derivative is obtained by: eliminating a reaction catalyst by filtration; and

subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

[2] Also, the substituted 5-alkylamino isoxazole derivative of the formula (V) can be prepared by the method shown in the following reaction scheme 2.



(In the reaction scheme 2, Hal represents a halogen atom, X, Y<sup>1</sup>, R<sup>1</sup>, and R<sup>2</sup> represent the same meanings as those defined above.)

That is, a 5-halogenoalkyl isoxazole derivative represented by the formula (VII) can be prepared by reacting an oxime derivative represented by the formula (II) with a propargyl amine derivative represented by the formula (VI) in an appropriate solvent in the presence of Hypolites.

Here, examples of Hypolites include: alkali metals

such as sodium hypochlorite, potassium hypochlorite, lithium hypochlorite, sodium hypobromite, and potassium hypobromite; calcium hypochlorite; magnesium hypochlorite; strontium hypochlorite; calcium hypobromite; and magnesium hypobromite.

5 Preferable examples thereof include alkali metal solutions of sodium hypochlorite, and potassium hypochlorite.

Preferable examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene,  
10 and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; esters such as methyl acetate, ethyl  
15 acetate, and propyl acetate; water; acetonitrile; dimethyl formamide; dimethyl sulfoxide; and N-methyl-2-pyrrolidone; and arbitrary mixed solvents thereof. More preferable examples thereof include: halogenated hydrocarbons such as chloroform and dichloromethane; and arbitrary mixed solvents  
20 thereof.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is  
25 generally completed in 30 minutes to 24 hours.

The compound of the formula (VII) which is a reaction intermediate is collected from a reaction mixture by a usual post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; washing with water; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

Subsequently, a substituted 5-aminoalkyl isoxazole represented by the formula (V) can be prepared by reacting the compound of the formula (VII) in an appropriate solvent by using an amine.

Examples of the amines used here include gaseous ammonia, an ammonia-dioxane solution, an ammonia-ethyl alcohol solution, an ammonia-methyl alcohol solution, an ammonia-2-propanol solution, and an aqueous ammonia solution.

Any solvent may be used for the reaction as long as the solvent does not inhibit the reaction: for example, hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; water; and arbitrary mixed solvents thereof.

More preferable examples thereof include halogenated



hydrocarbons such as chloroform and dichloromethane.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

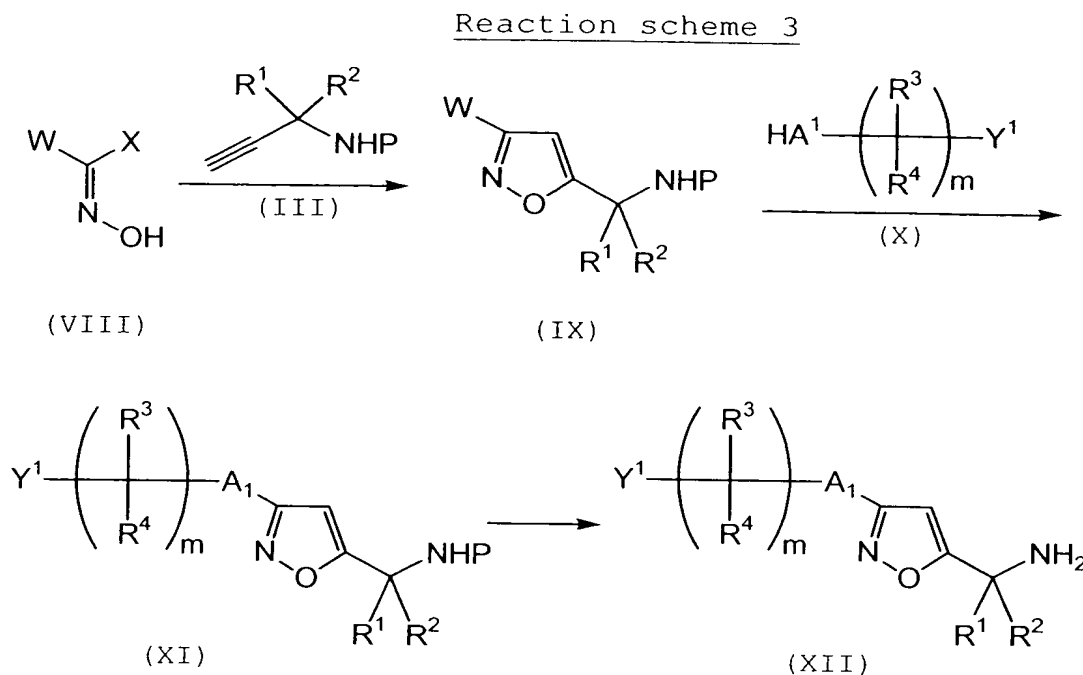
Although the reaction time depends on the reaction  
5 temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 1 week.

The compound of the formula (V) which is the reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by:  
10 making the reaction mixture alkali; then extracting with a solvent; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

15 The substituted 5-alkylamino isoxazole derivative of the formula (IV) described above can be synthesized by the methods described in JP 52-154525 A, JP 9-509951 A, Tetrahedron Letters, Vol. 34, No. 47, pp7509-7512, 1993, Tetrahedron Letters, Vol. 42 (2001) 1057-1060, Synthetic  
20 Communications, 22(13), 1939-1948 (1992), Tetrahedron Letters, Vol. 27, No. 27, pp3181-3182, 1986, Tetrahedron Letters, Vol. 33, No. 22, pp3113-3116, 1992, and the like, and the methods according thereto.

[3] A substituted 5-alkylamino isoxazole derivative  
25 represented by the formula (XII) can be prepared by the

method shown in the following reaction scheme 3.



(In the reaction scheme 3,  $\text{Y}^1$ ,  $\text{R}^1$ ,  $\text{R}^2$ ,  $\text{R}^3$ ,  $\text{R}^4$ , and  $m$  represent  
 5 the same meanings as those defined above,  $\text{A}^1$  represents an  
 oxygen atom, a sulfur atom, or an  $-\text{NR}^{12}-$  group (wherein  $\text{R}^{12}$   
 represents a hydrogen atom or a lower alkyl group),  $\text{Hal}$   
 represents a halogen atom,  $\text{P}$  represents a protecting group  
 of the amino group,  $\text{X}$  represents a halogen atom, and  $\text{W}$   
 10 represents a leaving group such as a halogen atom, or a  
 tosyl or mesyl group.)

That is, a protection body of a 3-phenylsulfonyl  
 isoxazole derivative represented by the formula (IX) can be  
 prepared by reacting halo oxime represented by the formula  
 15 (VIII) with the propargyl amine derivative represented by  
 the formula (III) by virtue of 1,3-dipolar addition in the  
 presence of a base. Subsequently, the protection body of a

5-alkylamino isoxazole derivative represented by the formula (XI) can be prepared by reacting a nucleophilic agent represented by the formula (X). A substituted alkylamino isoxazole derivative represented by the formula (XII) can be prepared by reacting the protection body of the alkylamino isoxazole derivative represented by the formula (XI) under an appropriate deprotection condition.

As the base used for the 1,3-dipolar addition reaction, as the base used in the 1st stage reaction for example, any type of base generally used for this type of reaction may be used without particular limitation. Examples thereof include: alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali earth metal hydroxides such as calcium hydroxide; alkali carbonates such as sodium carbonate and potassium carbonate; and organic bases such as triethylamine, trimethylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, 1,5-diazabicyclo[4.3.0]non-5-ene (DBN) and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU).

Any solvent may be used for the 1,3-dipolar addition reaction as long as the solvent does not inhibit the reaction: for example, hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform,

dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; water; and arbitrary mixed solvents thereof. Preferable examples thereof include halogenated hydrocarbons such as chloroform and  
5 dichloromethane.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is  
10 generally completed in 30 minutes to 1 week.

The compound of the formula (IX) which is a reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by: extracting with a solvent; and subsequently distilling off  
15 the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

As the base used in the nucleophilic addition reaction of the compound represented by the formula (X) with the  
20 compound represented by the formula (IX), any type of base generally used for this type of reaction may be used without particular limitation. Examples thereof include: alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali earth metal hydroxides such as calcium  
25 hydroxide; alkali carbonates such as sodium carbonate and

potassium carbonate; and organic bases such as triethylamine, trimethylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, 1,5-diazabicyclo[4.3.0]non-5-ene (DBN), and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU).

5 Any solvent may be used for the nucleophilic addition reaction of the compound represented by the formula (X) with the compound represented by the formula (IX) as long as the solvent does not inhibit the reaction including: for example, hydrocarbons such as hexane, pentane, and cyclohexane;  
10 aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-  
15 trichloroethane; water; and arbitrary mixed solvents thereof.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is  
20 generally completed in 30 minutes to 1 week.

The compound of the formula (XI) which is a reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by: extracting with a solvent; and subsequently distilling off  
25 the solvent. The resulting objective mixture can be also

purified by operations such as column chromatography and recrystallization if necessary.

Subsequently, a substituted 5-aminoalkyl isoxazole derivative represented by the formula (XII) can be prepared  
5 by reacting the compound of the formula (XI) in an appropriate solvent under an appropriate deprotection condition.

The deprotection condition used here depends on the protecting group. For example, when a t-butoxycarbonyl  
10 group is used as the protecting group, proton acid solutions such as an aqueous solution of hydrochloric acid are preferable, and when a benzyloxycarbonyl group is used as the protecting group, it is preferable to perform deprotection under a hydrogen flow in the presence of a  
15 catalyst such as palladium carbon or Raney Nickel.

Preferable examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and  
20 n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; esters such as methyl acetate, ethyl acetate, and propyl acetate; water; acetonitrile; dimethyl  
25 formamide; dimethyl sulfoxide; N-methyl-2-pyrrolidone; and

arbitrary mixed solvents thereof.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

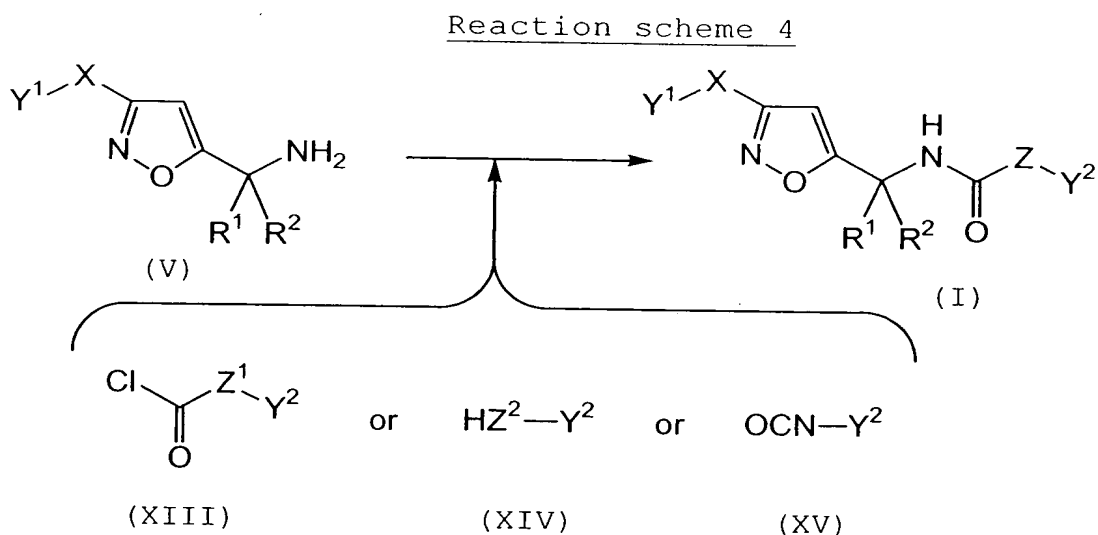
Although the reaction time depends on the reaction temperature, the reaction substrate, and the reaction reagent, the reaction is generally completed in 30 minutes to 1 week.

The substituted aminoalkyl isoxazole derivative of the formula (XII) is collected from the reaction mixture by the usual post-treatment. For example, in the case of the deprotection condition of the t-butoxycarbonyl group, the derivative is obtained by: making the reaction mixture alkali; then extracting with a solvent; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary. Also in the case of the deprotection condition of the benzyloxycarbonyl group, the derivative is obtained by: eliminating a reaction catalyst by filtration; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

Halo oxime represented by the formula (VIII) is the compound known in literatures, and for example, there are descriptions of the synthetic methods of the halo oxime and

the related reactions in Journal of American Chemical Society, 101, (5), 1319-1320 (1979), Journal of Organic Chemistry, 1988, 53, 5369-5371, Journal of Organic Chemistry, 1983, 46, 1796, Journal of Organic Chemistry, 1984, 49, 4595, 5 Journal of Organic Chemistry, 1987, 52, 2973, JP 11-504651 T, Synthetic Communications, 22(13), 1939-1948 (1992), Tetrahedron Letters, Vol. 27, No. 27, pp3181-3182, 1986, Tetrahedron Letters, Vol. 33, No. 22, pp3113-3116, (1992), and the like.

10 [4] The compound represented by the formula (I) can be prepared by the method shown in the following reaction scheme 4.



15 (In the reaction scheme, X, Y<sup>1</sup>, Y<sup>2</sup>, R<sup>1</sup>, R<sup>2</sup>, and Z represent the same meanings as those defined above, Z<sup>1</sup> represents Z in which B is any one but an NH group, and Z<sup>2</sup> represents Z which includes B.)

That is, the derivative represented by the formula (I)



can be prepared by: reacting a derivative, in which a carbonyl group of an acid chloride represented by the formula (XIII) or of a compound other than the compound of the formula (XIII) is activated, with the substituted 5-alkylamino isoxazole derivative (V) in an appropriate solvent, if necessary in the presence of an appropriate base.

Examples of the derivative in which a carboxyl group of a compound other than the compound of the formula (XIII) is activated used here include: an acid anhydride of the compound of the formula (XIII); a mixed acid anhydride in which the compound of the formula (XIII) is condensed with another acid, o-alkyl carbonate, or the like; and activated esters such as p-nitrophenyl ester, 2-tetrahydropyranyl ester, and 2-pyridyl ester.

The base used here is not particularly limited and may be any type of base generally used for this type of reaction. Examples of the base include: alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali earth metal hydroxides such as calcium hydroxide; alkali carbonates such as sodium carbonate, and potassium carbonate; and organic bases such as triethylamine, trimethylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, 1,5-diazabicyclo[4.3.0]non-5-ene (DBN), and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU). Of those, tertiary amines such as triethylamine, pyridine, and N-

methylnpiperidine are preferable.

Any solvent may be used for the reaction as long as the solvent does not inhibit the reaction: for example, hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; water; and arbitrary mixed solvents thereof.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 1 week.

The compound of the formula (I) which is the reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; extracting with a solvent; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

In this reaction, the substituted alkylamino isoxazole derivative represented by the formula (XII) may be used in

place of the substituted alkylamino isoxazole derivative represented by the formula (V).

Also, the compound of the formula (I) can be prepared by: making the substituted 5-alkylamino isoxazole derivative  
5 of the formula (V) act upon an active species obtained by reacting an alcohol derivative, thiol derivative, primary amino derivative, or secondary amino derivative denoted by the formula (XIV) in the presence of an appropriate solvent with an appropriate condensing agent, if necessary in the  
10 presence of an appropriate base.

Examples of the condensation agent used include N,N'-carbonyl diimidazole, 4-nitrophenoxy carbonyl chloride, phosgene, and triphosgene.

The base used here is not particularly limited and may  
15 be any type of base generally used for this type of reaction. Examples of the base include: alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali earth metal hydroxides such as calcium hydroxide; alkali carbonates such as sodium carbonate and potassium carbonate;  
20 and organic bases such as triethylamine, trimethylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, 1,5-diazabicyclo[4.3.0]non-5-ene (DBN), and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU). Of those, tertiary amines such as triethylamine, pyridine, and N-  
25 methylpiperidine are preferable.

Examples of the solvent used for the reaction include:  
hydrocarbons such as hexane, pentane, and cyclohexane;  
aromatic hydrocarbons such as benzene, toluene, and xylene;  
primary alcohols such as methanol, ethanol, and n-propanol;  
5 ethers such as tetrahydrofuran and dioxane; halogenated  
hydrocarbons such as chloroform, dichloromethane, carbon  
tetrachloride, 1,2-dichloroethane, and 1,1,2-  
trichloroethane; water; and arbitrary mixed solvents thereof.

The reaction temperature is generally between -80°C  
10 and 150°C, and preferably between -15°C and 50°C.

Although the reaction time depends on the reaction  
temperature and the reaction substrate, the reaction is  
generally completed in 30 minutes to 48 hours.

The compound of the formula (I) which is the reaction  
15 object is collected from the reaction mixture by the usual  
post-treatment. For example, the compound is obtained by:  
adding water to the reaction mixture; extracting with a  
solvent; and subsequently distilling off the solvent. The  
resulting objective mixture can be also purified by  
20 operations such as column chromatography and  
recrystallization if necessary.

In this reaction, the substituted alkylamino isoxazole  
derivative represented by the formula (XII) may be used in  
place of the substituted alkylamino isoxazole derivative  
25 represented by the formula (V).

Also, the compound of the formula (I) can be prepared by reacting an isocyanate derivative of the formula (XV) with the substituted 5-alkylamino isoxazole derivative of the formula (V).

5        Examples of the solvent used for the reaction include:  
hydrocarbons such as hexane, pentane, and cyclohexane;  
aromatic hydrocarbons such as benzene, toluene, and xylene;  
primary alcohols such as methanol, ethanol, and n-propanol;  
ethers such as tetrahydrofuran and dioxane; halogenated  
10 hydrocarbons such as chloroform, dichloromethane, carbon  
tetrachloride, 1,2-dichloroethane, and 1,1,2-  
trichloroethane; water; and arbitrary mixed solvents thereof.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

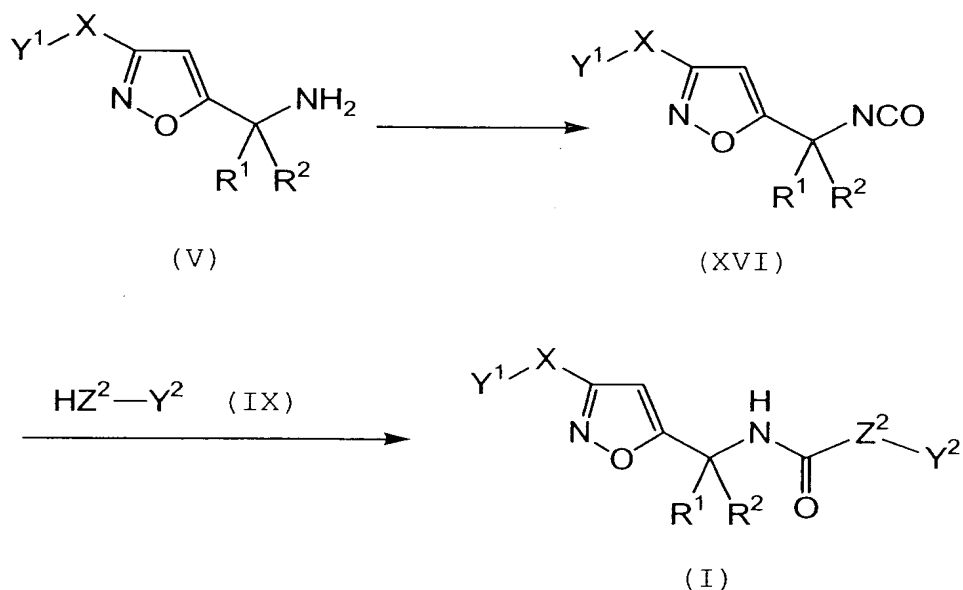
15        Although the reaction time depends on the reaction  
temperature and the reaction substrate, the reaction is  
generally completed in 30 minutes to 48 hours.

The compound of the formula (I) which is the reaction  
object is collected from the reaction mixture by the usual  
20 post-treatment. For example, the compound is obtained by:  
adding water to the reaction mixture; extracting with a  
solvent; and subsequently distilling off the solvent. The  
resulting objective mixture can be also purified by  
operations such as column chromatography and  
25 recrystallization if necessary.

In this reaction, the substituted alkylamino isoxazole derivative represented by the formula (XII) may be used in place of the substituted alkylamino isoxazole derivative represented by the formula (V).

5 [5] The derivative represented by the formula (I) can be also manufactured by the method shown in the following reaction scheme 5.

Reaction scheme 5



10 (In the reaction scheme,  $X$ ,  $Y^1$ ,  $Y^2$ ,  $R^1$ ,  $R^2$ , and  $Z^2$  represent the same meanings as those defined above.)

That is, the isocyanate derivative represented by the formula (XVI) can be prepared by using phosgene or triphosgene for the substituted 5-alkylamino isoxazole derivative (V) and reacting in an appropriate solvent, if  
 15 necessary in the presence of an appropriate base.

The base used here is not particularly limited and may

be any type of base generally used for this type of reaction. Examples of the base include: alkali metal hydroxides such as sodium hydroxide and potassium hydroxide; alkali earth metal hydroxides such as calcium hydroxide; alkali  
5 carbonates such as sodium carbonate and potassium carbonate; and organic bases such as triethylamine, trimethylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, 1,5-diazabicyclo[4.3.0]non-5-ene (DBN), and 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU). Of those, tertiary  
10 amines such as triethylamine, pyridine, and N-methylpiperidine are preferable.

Examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene;  
15 primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; water; and arbitrary mixed solvents thereof.

20 The reaction temperature is generally between -80°C and 150°C, and preferably between -15°C and 100°C.

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 48 hours.

25 The compound of the formula (XVI) which is the

reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; extracting with a solvent; and subsequently distilling off  
5 the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

In this reaction, the substituted alkylamino isoxazole derivative represented by the formula (XII) may be used in  
10 place of the substituted alkylamino isoxazole derivative represented by the formula (V).

Subsequently, the compound of the formula (I) can be also manufactured by reacting an alcohol derivative, thiol derivative, primary amino derivative, or secondary amino  
15 derivative represented by the formula (IX) with the isocyanate derivative (XVI) in the presence of an appropriate solvent.

Examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane;  
20 aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane; halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-  
25 trichloroethane; water; and arbitrary mixed solvents thereof.



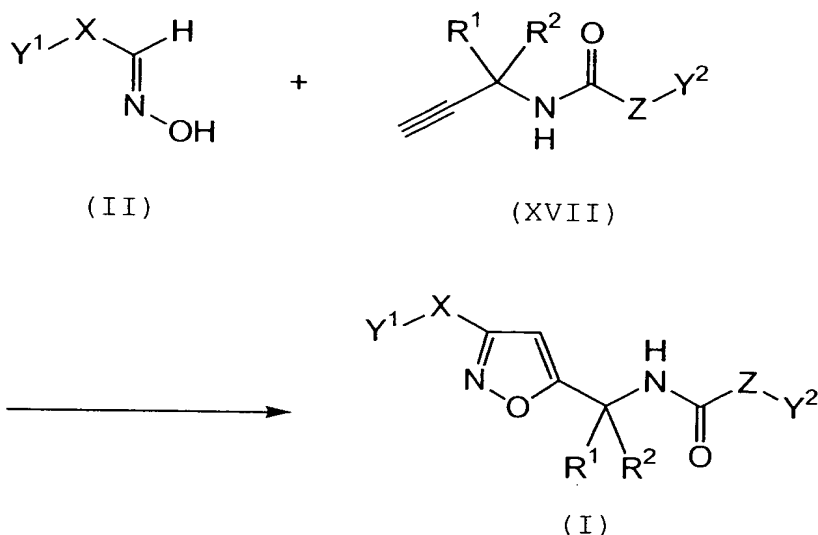
The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 48 hours.

The compound of the formula (I) which is the reaction object is collected from the reaction mixture by the usual post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; extracting with a solvent; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

[6] The substituted 5-alkylamino isoxazole derivative represented by the formula (I) can be also manufactured by the method shown in the following reaction scheme 6.

Reaction scheme 6



(In the reaction scheme,  $X$ ,  $Y^1$ ,  $Y^2$ ,  $R^1$ ,  $R^2$ , and  $Z$  represent the same meanings as those defined above.)

That is, the derivative represented by the formula (I) can be prepared by reacting the oxime derivative represented  
5 by the formula (II) with a propargyl amine derivative represented by the formula (XVII) in an appropriate solvent in the presence of Hypolites.

Here, examples of Hypolites include: alkali metals such as sodium hypochlorite, potassium hypochlorite, lithium  
10 hypochlorite, sodium hypobromite, and potassium hypobromite; calcium hypochlorite; magnesium hypochlorite; strontium hypochlorite; calcium hypobromite; and magnesium hypobromite. Preferable examples thereof include alkali metal solutions of sodium hypochlorite and potassium hypochlorite.

15 Preferable examples of the solvent used for the reaction include: hydrocarbons such as hexane, pentane, and cyclohexane; aromatic hydrocarbons such as benzene, toluene, and xylene; primary alcohols such as methanol, ethanol, and n-propanol; ethers such as tetrahydrofuran and dioxane;  
20 halogenated hydrocarbons such as chloroform, dichloromethane, carbon tetrachloride, 1,2-dichloroethane, and 1,1,2-trichloroethane; esters such as methyl acetate, ethyl acetate, and propyl acetate; water; acetonitrile; dimethyl formamide; dimethyl sulfoxide; N-methyl-2-pyrrolidone; and  
25 arbitrary mixed solvents thereof. More preferable examples

thereof include halogenated hydrocarbons such as chloroform and dichloromethane.

The reaction temperature is generally between  $-80^{\circ}\text{C}$  and  $150^{\circ}\text{C}$ , and preferably between  $-15^{\circ}\text{C}$  and  $50^{\circ}\text{C}$ .

5        Although the reaction time depends on the reaction temperature and the reaction substrate, the reaction is generally completed in 30 minutes to 24 hours.

10        The compound of the formula (I) is collected from a reaction mixture by a usual post-treatment. For example, the compound is obtained by: adding water to the reaction mixture; washing with water; and subsequently distilling off the solvent. The resulting objective mixture can be also purified by operations such as column chromatography and recrystallization if necessary.

15        Aldoximes of the compound (II) can be prepared from corresponding aldehydes using a base such as hydroxylamine hydrochloride or sodium acetate in an alcohol solvent such as ethanol at ambient temperature or under the condition of reflux with heating.

20        Also, the propargyl amine derivative of the formula (XVII) can be prepared from propargyl amine using chloroformates, isocyanates, acid chlorides, or the like.

[Application as agri-horticultural fungicides]

25        The compound represented by the formula (I) or (IA)

has an action of exterminating noxious organisms and can be used as a agri-horticultural fungicide.

Plant diseases toward which the compound of the present invention have an excellent fungicidal effect  
5 include:

*Pyricularia oryzae*;

*Rhizoctonia solani*;

*Gibberella fujikuroi*;

*Erysiphe graminis* f.sp. *tritici*;

10 *Erysiphe graminis* f.sp. *hordei*;

*Pseudocercospora herpotrichoides*;

*Puccinia graminis*;

*Colletotrichum graminicola*;

*Septoria tritici*;

15 *Phynchosporium secalis* f.sp. *hordei*;

*Phytophthora infestans*;

*Alternaria solani*;

*Colletotrichum atramentarium*;

*Thanatephorus cucumeris*;

20 *Botrytis cinerea*;

*Erysiphe pisi*;

*Cercospora canescens*;

*Sclerotinia sclerotiorum*;

*Colletotrichum phaseolorum*;

25 *Cercospora canescens*;

- Sclerotinia sclerotiorum*;  
*Colletotrichum lindemuthianum*;  
*Botrytis cinerea*;  
*Colletotrichum truncatum*;  
5 *Cercospora kikuchii*;  
*Phakopsora pachyrhizi*;  
*Gloeosporium conjac*;  
*Septoria perillae*;  
*Colletotrichum theae-sinensis*;  
10 *Cercospora beticola*;  
*Colletotrichum spinaciae*;  
*Peronospora effusa*;  
*Alternaria brassicae*;  
*Alternaria brassicicola*;  
15 *Sclerotinia sclerotiorum*;  
*Botrytis cinerea*;  
*Colletotrichum higginsianum*;  
*Sclerotinia sclerotiorum*;  
*Alternaria brassicae*, *Alternaria brassicicola*;  
20 *Alternaria cucumerina*;  
*Botrytis cinerea*;  
*Pseudoperonospora cubensis*;  
*Sphaerotheca fuliginea*;  
*Phytophthora melonis*;  
25 *Corynespora cassiicola*;

- Sclerotinia sclerotiorum*;
- Alternaria cucumerina*;
- Colletotrichum lagenarium*;
- Fusarium oxysporum* f.sp. *cucumerinum*;
- 5 *Pythium cucurbitacearum*, *Rhizoctonia solani*;
- Botrytis cinerea*;
- Phomopsis* sp.;
- Pseudoperonospora cubensis*;
- Sphaerotheca fuliginea*;
- 10 *Phytophthora cryptogea*;
- Sclerotinia sclerotiorum*;
- Colletotrichum orbiculare*;
- Pseudoperonospora cubensis*;
- Corynespora cassiicola*;
- 15 *Sclerotinia sclerotiorum*;
- Fusarium oxysporum* f.sp. *melonis*;
- Pseudoperonospora cubensis*;
- Fusarium oxysporum*;
- Sclerotinia sclerotiorum*;
- 20 *Puccinia cnici-oleracei*;
- Gloeosporium chrysanthemi*, *Gloeosporium carthami*;
- Fuŝarium oxysporum*;
- Erysiphe heraclei*;
- Sclerotinia intermedia*, *Sclerotinia sclerotiorum*;
- 25 *Alternaria dauci*;

- Alternaria radicina*;
- Fusarium oxysporum*;
- Oidiopsis sicula*;
- Phytophthora capsici*;
- 5 *Sclerotinia sclerotiorum*;
- Colletotrichum gloeosporioides*;
- Botrytis cinerea*;
- Fusarium oxysporum* f.sp. *lycopersici*;
- Oidiopsis sicula*;
- 10 *Phytophthora infestans*;
- Sclerotinia sclerotiorum*;
- Botrytis cinerea*;
- Fulvia fulva*;
- Phytophthora infestans*;
- 15 *Thanatephorus cucumeris*;
- Phytophthora capsici*;
- Sclerotinia sclerotiorum*;
- Alternaria solani*;
- Fusarium oxysporum*;
- 20 *Verticillium dahliae*;
- Sphaerotheca humuli*;
- Phytophthora nicotianae* var. *parasitica*;
- Pythium ultimum* var. *ultimum*;
- Sclerotinia sclerotiorum*;
- 25 *Alternaria alternata*;

- Mycosphaerella fragariae*;  
*Colletotrichum acutatum*, *Glomerella cingulata*;  
*Botrytis cinerea*;  
*Cercospora asparagi*;  
5 *Phomopsis asparagi*;  
*Puccinia asparagi-lucidi*;  
*Colletotrichum gloeosporioides*;  
*Botrytis cinerea*;  
*Phytophthora nicotianae*;  
10 *Cladosporium allii-cepae*;  
*Fusarium oxysporum* f.sp. *cepae*;  
*Sclerotinia sclerotiorum*;  
*Septoria alliacea*;  
*Alternaria porri*;  
15 *Puccinia allii*;  
*Puccinia allii*;  
*Botrytis squamosa*;  
*Phytophthora porri*;  
*Colletotrichum circinans*;  
20 *Alternaria* sp.;  
*Botrytis cinerea*;  
*Botrytis allii*;  
*Pleospora herbarum*;  
*Peronospora destructor*;  
25 *Puccinia allii*;



- Botrytis byssoidea*;  
*Phytophthora nicotianae*;  
*Fusarium oxysporum*;  
*Mycosphaerella allicina*;  
5 *Septoria alliacea*;  
*Alternaria porri*;  
*Sclerotinia allii*;  
*Puccinia allii*;  
*Botrytis squamosa*;  
10 *Phytophthora porri*;  
*Colletotrichum circinans*;  
*Peronospora destructor*;  
*Colletotrichum gloeosporioides*;  
*Phyllactinia kakicola*;  
15 *Colletotrichum* ssp., *Glomerella* sp.;  
*Cercospora kakivora*;  
*Cercospora kaki*;  
*Macrophoma kaki*;  
*Fusicladium levieri*;  
20 *Phomopsis kakivora*;  
*Pseudocercospora fuliginosa*;  
*Physalospora kaki*;  
*Aureobasidium pullulans*, *Capnophaeum fuliginodes*,  
*Cladosporium herbarum*, *Microxyphium* sp., *Scorias communis*,  
25 *Tripospermum juglandis*;

- Zygophiala jamaicensis*;  
*Gloeosporium kaki*;  
*Botrytis cinerea*;  
*Pestalotia diospyri*;  
5 *Glomerella cingulata*;  
*Mycosphaerella nawae*;  
*Podosphaera tridactyla*, *Sphaerotheca pannosa*;  
*Botryosphaeria dothidea*;  
*Sclerotinia sclerotiorum*;  
10 *Cladosporium carpophilum*;  
*Leucotelium pruni-persicae*;  
*Rosellinia necatrix*;  
*Fusarium lateritium*;  
*Pseudocercospora circumscissa*;  
15 *Sphaceloma pruni-domesticae*;  
*Glomerella cingulata*;  
*Botrytis cinerea*;  
*Monilinia fructicola*, *Monilinia laxa*;  
*Glomerella mume*;  
20 *Sclerotinia sclerotiorum*;  
*Podosphaera tridactyla*;  
*Rhizopus nigricans*;  
*Glomerella cingulata*;  
*Botrytis cinerea*;  
25 *Monilinia fructicola*;

- Phyllactinia mali*;
- Phytophthora cactorum*, *Phytophthora syringae*;
- Venturia pirina*;
- Colletotrichum gloeosporioides*;
- 5 *Gymnosporangium asiaticum*;
- Phyllactinia pyri*;
- Phytophthora cactorum*, *Phytophthora syringae*;
- Venturia nashicola*;
- Alternaria kikuchiana*;
- 10 *Leptothyrium pomi*;
- Glomerella cingulata*;
- Botrytis cinerea*;
- Monilinia fructigena*;
- Colletotrichum gloeosporioides*;
- 15 *Physalospora piricola*;
- Fusarium oxysporum*, *Gibberella zeae*;
- Podosphaera tridactyla*, *Sphaerotheca pannosa*;
- Sclerotinia sclerotiorum*;
- Rhizopus nigricans*;
- 20 *Cladosporium carpophilum*;
- Leucotelium pruni-persicae*;
- Stenella* sp.;
- Pseudocercospora circumscissa*, *Phyllosticta persicae*;
- Gloeosporium laeticolor*;
- 25 *Botrytis cinerea*;

*Monilinia fructicola*;  
*Phomopsis* sp.;  
*Gymnosporangium yamadae*;  
*Podosphaera leucotricha*;  
5      *Phytophthora cactorum*,      *Phytophthora cambivora*,  
*Phytophthora syringae*;  
*Diplocarpon mali*;  
*Cristulariella moricola*;  
*Venturia inaequalis*;  
10      *Zygophiala jamaicensis*;  
*Glomerella cingulata*;  
*Botrytis cinerea*;  
*Alternaria alternata*;  
*Uncinula necator*;  
15      *Sclerotinia sclerotiorum*;  
*Glomerella cingulata*;  
*Pseudocercospora vitis*;  
*Briosia ampelophaga*;  
*Elsinoe ampelina*;  
20      *Phyllosticta ampellicida*;  
*Phomopsis viticola*;  
*Plasmopara viticola*;  
*Glomerella cingulata*;  
*Capnodium salicinum*;  
25      *Morenoella quercina*;

*Microsphaera alphitoides;*

*Monochaetia monochaeta;*

*Botrytis cinerea;*

*Phytophthora citrophthora;*

5 *Diaporthe citri;*

*Mycosphaerella citri, Mycosphaerella horii;*

*Elsinoe fawcettii;*

*Mycosphaerella pinodes;*

*Colletotrichum gloeosporioides;* and

10 *Colletotrichum gloeosporioides.*

Application of the compound by the invention can be performed by treating plants with an active ingredient by spraying, scattering, applying, or the like, or by treating seeds of the plants, soil surrounding the plants or soil in  
15 which the seeds are sowed, and rice paddies and water of hydroponics with the active ingredient. The application can be performed before or after the plant is infected with pathogenic fungi.

The compound of the invention can be used by combining  
20 and formulating carriers, surfactants, adjuncts, and the like commonly used in agrichemical formulations.

The aspects of the formulations include dosage forms such as suspension agents, oil agents, emulsions, liquid agents, water soluble agents, water-dispersible powder,  
25 flowable agents, paste agents, granules, powder, tablets,

granular water-dispersible powder, fumigants, aerosol agents, foaming agents, microcapsule agents, jumbo agents, coating agents for seeds, and ULV (cold mists or warm mists) agents. Such aspects can be prepared by the usual method of mixing  
5 at least one compound of the invention with carriers such as appropriate solid carrier, liquid carrier, and liquid gaseous carrier, and if desired, appropriate adjuncts (e.g., surfactants, spreaders, dispersants, stabilizers) for improving dispersibility and other natures of the active  
10 ingredient.

Examples of the solid carrier include: vegetable materials (such as crystalline cellulose, starch, wood meal, cork, and coffee shells); fibrous materials; artificial plastic powders; clays (such as kaolin, bentonite, white  
15 clay, diatomite, synthetic hydroxylated silicon, fubasami clay, and acid clay); talc and inorganic products (such as vermiculite, montmorillonite, pumica, sulfur powder, apatite, mica, sericite, ground quartz, active carbon, and calcium carbonate); high-molecular compounds (such as  
20 polyvinylchloride and petroleum resin); and chemical fertilizers (such as ammonium sulfate, ammonium phosphate, ammonium nitrate, ammonium chloride, calcium chloride, and urea).

Examples of the liquid carrier and diluent include:  
25 water; alcohols (such as methanol, ethanol, isopropyl

alcohol, cyclohexanol, and aluminium alcohol); ketones (such as acetone, methyl ethyl ketone, and cyclohexanone); ethers (such as ethylcellosolve, butylcellosolve, and dioxane); aromatic hydrocarbons (such as benzene, toluene, xylene, ethylbenzene, and methylnaphthalene); aliphatic hydrocarbons (such as kerosine, isoparaffin, and normal paraffin); esters (such as isopropyl acetate and benzyl acetate); nitriles; amides (such as N,N-dimethylformamide and dimethylsulfoxide); and halogenated hydrocarbons (such as chlorobenzene and trichloroethylene).

Examples of the gaseous carrier, in other words the propellant, include carbon dioxide gas, butane gas, and fluorocarbon.

Examples of the surfactant include: alkyl sulfate; alkyl sulfonate; alkyl arylsulfonate; alkyl arylates and polyoxyethylene compounds thereof; polyethylene glycolether; polyalcohol esters; and sugar alcohol derivatives.

Examples of other adjuncts for the preparation include: binders and dispersants such as casein, gelatin, polysaccharides (such as starch, gum arabic, cellulose derivatives, and alginic acid), lignin derivatives, bentonite, synthetic water-soluble high molecules (such as polyvinyl alcohol, polyvinyl pyrrolidone, and polyacrylic acid); and stabilizers such as PAP (acidic phosphoric isopropyl), BHT (2,6-di-tert-butyl-4-methylphenol), BHA (2-

tert-butyl-4-methoxyphenol), vegetable oil, molasses, agar, mineral oil, fatty acid, and fatty ester.

Generally, each of the above preparations contains at least one compound (I) of the invention at 0.1 to 95% by weight, preferably at 1 to 80% by weight. Those preparations can be treated alone or in dilution for the plants by spraying, scattering, applying, or the like, or for seeds of the plants, soil surrounding the plants or soil in which the seeds are sowed, and rice paddies and water of hydroponics. When the compound of the invention is sprayed, scattered, or applied onto the plants, the compound is used at a concentration of about 1 to 50,000 ppm, preferably about 50 to 1,000 ppm. When treated for the soil surrounding the plants or water of rice paddies, the compound is used at an amount of about 1 g to 5 kg/hectare, preferably about 2 g to 100 g/hectare. When treated for the soil in which the seeds are sowed, the compound is typically used at an amount of about 0.1 g to 1 kg/m<sup>2</sup>, preferably 0.5 g to 500 g/m<sup>2</sup>.

The compound of the invention can be formulated by combining with one or more of the other agrichemical ingredients (fungicide, bactericide, insecticide, acaridide, herbicide, plant growth regulator, etc.) for expanding effects. Examples of such agrichemicals are shown below, but the examples of agrichemicals to be combined are not



limited thereto.

Examples of the combinable fungicide include acibenzolar-s-methyl, azoxystrobin, isoprothiolane, ipconazole, iprodione, iprovalicarb, iminoctadine acetate, iminoctadine albesilate, epoxyconazole, basic copper chloride, oxadixyl, kasugamycin, carpropamid, quinoxifen, captan, kresoxim-methyl, chlorothalonil, diazofamid, diethofencarb, diclocymet, diclomedin, dithianon, zineb, simeconazole, dimethirimol, cyproconazole, cymoxanil, silthiofam, streptomycin, spiroxamine, sulfenic acid, thiabenzazole, thiophanete methyl, thiuram, thifluzamide, tecloftalam, tebuconazole, triadimefon, triazine, tricyclazole, triflumizole, trifloxystrobin, triforine, triclofos methyl, copper nonylphenol sulfonate, paclobutrazol, validamycin, picoxystrobin, hydroxyisoxazole, pyraclostrobin, pyroquilon, famoxadone, fenarimol, fenoxanil, ferimzone, fenhexamid, fthalide, blasticidin S, furametpyr, fluazinam, fluquinconazole, fludioxonil, flutolanil, prochloraz, procymidone, propiconazole, probenazole, pefurazoate, benomyl, pencycuron, fosetyl, polyoxine, manzeb, metalaxyl, metominostrobin, mepanipyrim, mepronil, organic copper, organic nickel, edifenphos, and iprobenfos.

Examples of the combinable bactericide include tecloftalam and oxolinic acid.

Examples of the combinable insecticide include

acetamiprid, acephate, isoxathion, imidacloprid,  
ethiofencarb, ethylthiometon, ethofenprox, oxamyl, cartap,  
carbosulfan, clothianidin, chromafenozide, chlorpyrifos  
methyl, cycloprothrin, dinotefuran, dimethylvinphos,  
5 dimethoate, cyfluthrin, diprofezine, silafluofen, spinosad,  
sulprofos, diazinon, thiacloprid, thiomethoxam, thiodicarb,  
thiocyclam, thiometon, tebufenozide, nitenpyram, vamidothion,  
pymetrozine, pyraclofos, pyridaphenthion, fipronil, fopronil,  
buprofezin, furathiocarb, flucythrinate, fluvalinate,  
10 propaphos, permethrin, bensultap, benthiazole, benfuracarb,  
malathion, methasulfocarb, monocrotophos, malathion,  
fenobucarb, tetrachlorvinphos, trichlorophon, EPN,  
iprobenfos, fenitrothion, isoprocab, carbaryl, phenthoate,  
propoxur, and XMC.

15 Examples of the combinable acaricides include  
etoxazole, kelthane, fenbutatin oxide, tebufenpyrad,  
halfenprox, bifenazate, pyrimidifen, pirimiphos methyl,  
fenpropathrin, fluvalinate, phosalone, milbemectin,  
propargite, and dichlorvos.

20 Examples of the combinable herbicide include ioxynil,  
azimsulfuron, asulam, atrazine, anilofos, alachlor, imazamox  
ammonium, imazosulfuron, indanofan, esprocarb,  
ethoxysulfuron, etobenzanid, chlorate, oxadiclefomefon,  
cafenstrole, quizalofop ethyl, cumyluron, glyphosate  
25 ammonium, glyphosate isopropylamine, glyphosate trimesium,

glyphosate sodium, glyphosinate, clethodim, clomeprop, cyanazine, cyclosulfamuron, diquat, cyhalofop butyl, cyhalofop methyl, diflufenican, dimethametryn, dimethenamid, simetryn, dimepiperate, cinmethylin, lime nitrogen, 5 sethoxydim, dymron, thifensulfuron methyl, thenylchlor, tepraloxydim, trifluralin, naproanilide, paraquat, halosulfuron methyl, bialaphos, bispyribac sodium, bifenox, piperophos, pyrazoxyfen, pyrazosulfuron ethyl, pyrazolate, pyraflufenethyl, pyridate, pyributicarb, pyriminobac methyl, 10 fenoxapropethyl, fentrazamide, butachlor, butamifos, fluazifop, pretilachlor, prometryn, bromobutide, pelargonic acid, bensulfuron methyl, benzobicyclon, benzofenap, bentazone, benthocarb, pendimethalin, pentoxazone, benfuresate, methyl isocyanate, metolachlor, metribuzin, 15 mephenacet, molinate, linuron, lenacil, quinoclamine, simazine, dichlobenil, diuron, propanil, chlorpropham, MCPA ethyl, MCPA thioethyl, MCPA sodium, MCPB, bensulide, and 2,4-PA.

Examples of the combinable plant growth regulator 20 include inabenfide, indoleacetic acid, uniconazole, calcium peroxide, nicotinamide, paclobutrazol, prohexadione calcium, and benzylaminopurin.

#### **BEST MODE FOR CARRYING OUT THE INVENTION**

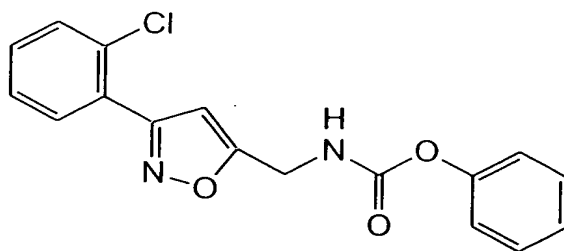
25 Next, the present invention is illustrated by citing

preparation examples and evaluation examples of the compound of the invention, but the invention is not limited thereto.

In the examples, data on  $^1\text{H-NMR}$  were represented as  $\delta$  values measured by an R-1900 type spectrometer (90 MHz) supplied from Hitachi Ltd., using tetramethylsilane as an internal standard and  $\text{CDCl}_3$  as a solvent. However, for the compound which is difficult to dissolve in  $\text{CDCl}_3$ , the measurement was performed by adding an appropriate amount of  $\text{DMSO-d}_6$ .

10

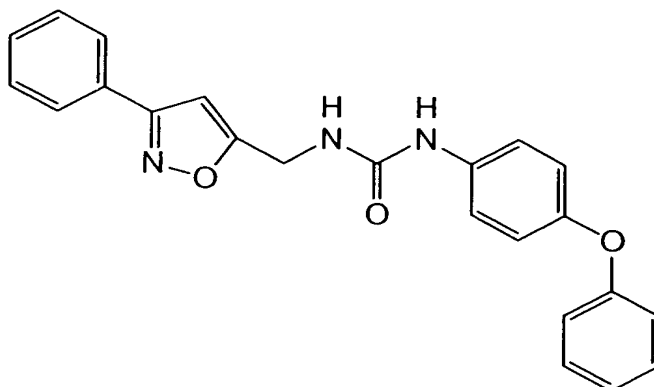
Example 1: Preparation of phenyl {[3-(2-chlorophenyl)-5-isoxazolyl]methyl}carbamate (Compound Number I-28)



To a solution of 0.51 g of 5-aminomethyl-3-(2-chlorophenyl)isoxazole and 0.38 g of diisopropylethylamine in 30 ml of dichloromethane, 0.46 g of phenyl chloroformate was added at  $0^\circ\text{C}$ , the temperature was raised to room temperature, and subsequently the solution was stirred for 5 hours. After the completion of the reaction, the solution was concentrated under reduced pressure, and subsequently purified on silica gel column chromatography to yield 0.75 g as a colorless oil having the following physical property.

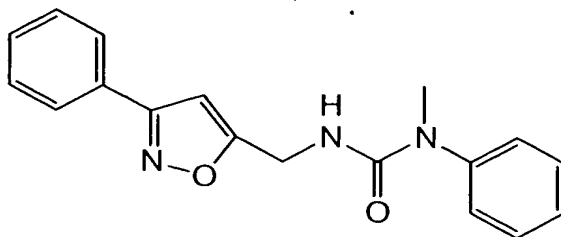
NMR :  $\delta$  4.52 (2H,d), 5.49-5.88 (1H,brs), 6.48 (1H,s), 6.99-7.80 (9H,m)

Example 2: Preparation of 3-[(3-phenyl-5-isoxazolyl)methyl]-  
5 1-(4-phenoxyphenyl)urea (Compound Number II-42)



To a solution of 0.82 g of 5-aminomethyl-3-phenylisoxazole in 40 ml of ethanol, 1.00 g of p-phenoxyphenylisocyanate was added at room temperature, and  
10 then the whole was stirred overnight. The reaction solution was concentrated under reduced pressure, and subsequently a precipitated crystal was filtrated and washed with isopropylether to yield 1.60 g as a colorless crystal having the following physical property.  
15 Melting point: 179 to 181°C.

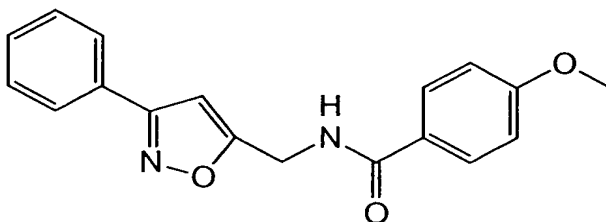
Example 3: Preparation of 3-[(3-phenyl-5-isoxazolyl)methyl]-  
1-methyl-1-phenyl urea (Compound Number II-12)



To a solution of 1.00 g of 5-aminomethyl-3-phenylisoxazole and 0.89 g of diisopropylethylamine in 20 ml of dichloromethane, 0.97 g of N-methyl N-phenylcarbamoylchloride was added at 0°C, the temperature was raised to room temperature, and subsequently the solution was stirred overnight. After the completion of the reaction, the solution was concentrated under reduced pressure, and subsequently purified on silica gel column chromatography to yield 1.48 g as a colorless crystal having the following physical property.

Melting point: 114 to 117°C.

Example 4: Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-methoxy benzamide (Compound Number III-111)

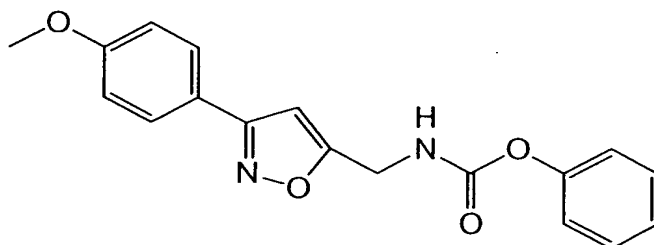


To a solution of 0.60 g of 5-aminomethyl-3-phenylisoxazole and 0.53 g of diisopropylethylamine in 30 ml of dichloromethane, 0.59 g of p-methoxybenzoylchloride was

added at 0°C, the temperature was raised to room temperature, and subsequently the solution was stirred for overnight. After the completion of the reaction, the solution was concentrated under reduced pressure, and subsequently  
5 purified on silica gel column chromatography to yield 0.90 g as a colorless crystal having the following physical property.

Melting point: 182 to 184°C.

- 10 Example 5: Preparation of phenyl {[3-(4-methoxyphenyl)-5-isoxazolyl]methyl}carbamate (Compound Number I-26)

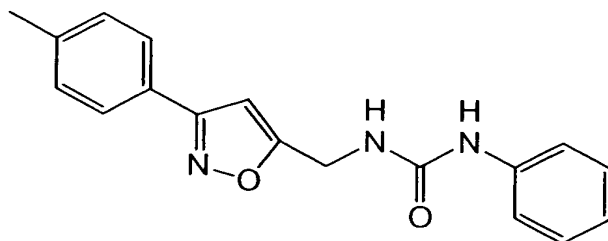


- To a solution of 2.00 g of 4-methoxy benzaldoxime and 2.78 g of phenyl 2-propynyl carbamate in 50 ml of  
15 dichloromethane, 11.8 g of 10% sodium hypochlorite aqueous solution was dripped over 30 minutes at 0°C. The temperature was raised to room temperature, and then the solution was stirred for 5 hours. Water was added, the solution was extracted twice with ethyl acetate, and an extract was  
20 washed with a saturated aqueous solution of sodium sulfite. The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 2.25

g as a colorless powder having the following physical property.

Melting point: 102 to 105°C.

5 Example 6: Preparation of 3-([3-(4-tolyl)-5-isoxazolyl]methyl)-1-phenyl urea (Compound Number II-2)



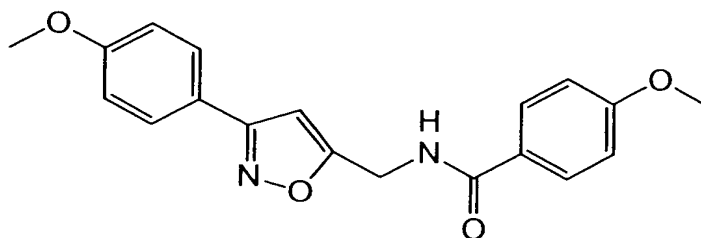
To a solution of 4.66 g of 4-methyl benzaldoxime and 1.50 g of 3-(2-propynyl)-1-phenyl urea in 50 ml of  
10 dichloromethane, 36.6 g of 7.0% sodium hypochlorite aqueous solution was dripped over 1 hour at 0°C. The temperature was raised to room temperature, and then the solution was stirred for 7 hours. Water was added, the solution was extracted twice with ethyl acetate, and an extract was  
15 washed with a saturated aqueous solution of sodium sulfite. The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 1.87 g as a colorless powder having the following physical property.

20 Melting point: 176 to 178°C.

Example 7: Preparation of N-([3-(4-methoxyphenyl)-5-



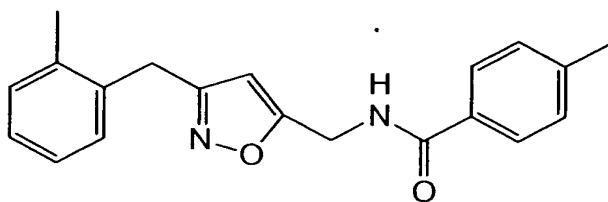
isoxazolyl)methyl}-4-methoxy benzamide (Compound Number III-18)



To a solution of 4.90 g of 4-methoxy benzaldoxime and  
5 1.50 g of N-(2-propynyl)-1-p-methoxy benzamide in 50 ml of  
dichloromethane, 37.7 g of 6.4% sodium hypochlorite aqueous  
solution was dripped over 1 hour at 0°C. The temperature was  
raised to room temperature, and then the solution was  
stirred overnight. Water was added, the solution was  
10 extracted twice with ethyl acetate, and an extract was  
washed with a saturated aqueous solution of sodium sulfite.  
The extract was dried on anhydrous magnesium sulfate, and  
subsequently purified on column chromatography to yield 2.36  
g as a colorless powder having the following physical  
15 property.

Melting point: 144 to 155°C.

Example 8: Preparation of N-([3-(2-methylbenzyl)-5-  
isoxazolyl)methyl]-4-methyl benzamide (Compound Number IV-  
20 12)

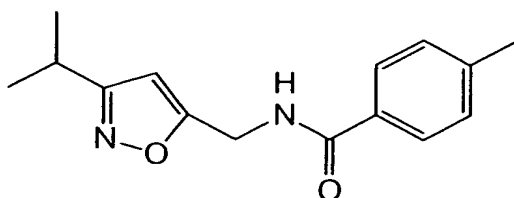


To a solution of 7.75 g of 2-methyl benzylaldoxime and 1.50 g of N-(2-propynyl)-1-p-methyl benzamide in 70 ml of dichloromethane, 87.9 g of 4.4% sodium hypochlorite aqueous solution was dripped over 30 minutes at 0°C. The temperature was raised to room temperature, and then the solution was stirred for 2 hours. Water was added, the solution was extracted twice with ethyl acetate, and an extract was washed with a saturated aqueous solution of sodium sulfite.

The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 2.12 g as a colorless oil having the following physical property.

NMR :  $\delta$  2.27(3H,s), 2.36(3H,s), 3.93(2H,s), 4.62(2H,d), 5.83(1H,s), 6.48-6.78(1H,brs), 7.13(4H,s), 7.20(2H,d), 7.67(2H,d)

Example 9: Preparation of N-[(3-isopropyl-5-isoxazolyl)methyl]-4-methyl benzamide (Compound Number III-5)



To a solution of 7.54 g of isopropylaldoxime and 1.50 g of N-(2-propynyl)-1-p-methyl benzamide in 100 ml of

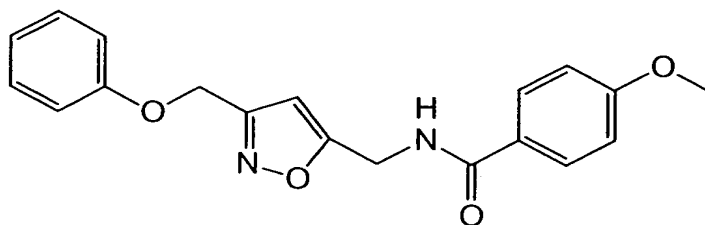
dichloromethane, 161.2 g of 4.0% sodium hypochlorite aqueous solution was dripped over 30 minutes at 0°C. The temperature was raised to room temperature, and then the solution was stirred overnight. Water was added, the solution was  
5 extracted twice with ethyl acetate, and an extract was washed with a saturated aqueous solution of sodium sulfite. The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 0.56 g of a colorless oil having the following physical property.

10 Melting point: 107 to 116°C.

NMR :  $\delta$  1.25(6H,d), 2.38(3H,s), 2.78-3.24(1H,m), 4.69(2H,d), 6.08(1H,s), 6.85(1H,br), 7.21(2H,d), 7.71(2H,d)

Example 10: Preparation of N-[(3-phenyloxymethyl-5-isoxazolyl)methyl]-4-methoxy benzamide (Compound Number IV-

15 45)



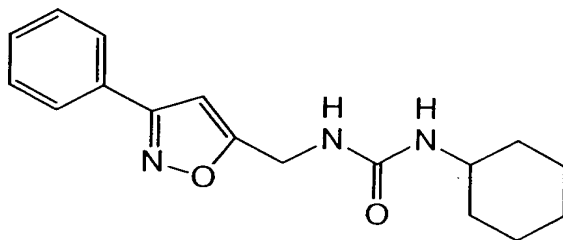
To a solution of 5.99 g of phenyloxymethylaldoxime and 1.50 g of N-(2-propynyl)-1-p-methoxy benzamide, 65.6 g of 4.5% sodium hypochlorite aqueous solution was dripped over  
20 30 minutes at 0°C. The temperature was raised to room temperature, and then the solution was stirred overnight. Water was added, the solution was extracted twice with ethyl

acetate, and an extract was washed with a saturated aqueous solution of sodium sulfite. The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 1.00 g as a colorless paste  
5 having the following physical property.

NMR :  $\delta$  3.82(3H,s), 4.72(2H,d), 5.12(2H,s), 6.34(1H,s), 6.41-6.65(1H,brs), 6.75-7.88(9H,m)

Example 11: Preparation of 3-[(3-phenyl-5-isoxazolyl)methyl]-1-cyclohexyl urea (Compound Number II-7)

10

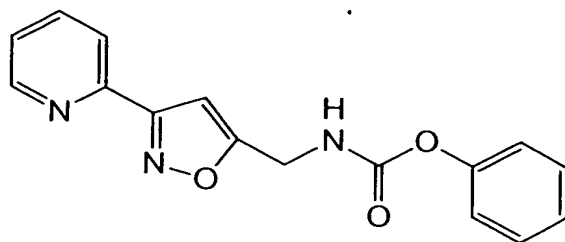


To a solution of 1.00 g of 5-aminomethyl-3-phenylisoxazole in 30 ml of ethanol, 0.79 g of cyclohexylisocyanate was added at room temperature, and then the whole was stirred overnight. The reaction solution was  
15 concentrated under reduced pressure, and subsequently a precipitated crystal was filtrated and washed with isopropylether to yield 1.65 g as a colorless crystal having the following physical property.

Melting point: 155 to 158°C.

20

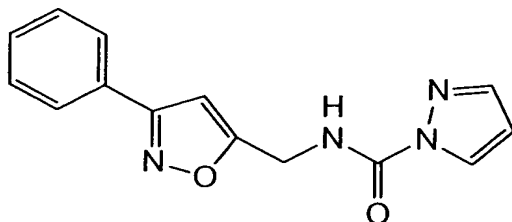
Example 12: Preparation of phenyl {[3-(2-pyridyl)-5-isoxazolyl)methyl}carbamate (Compound Number I-31)



To a solution of 1.00 g of 5-aminomethyl-3-(2-pyridyl)isoxazole and 0.89 g of diisopropylethylamine in 30 ml of dichloromethane, 0.89 g of phenyl chloroformate was added at 0°C, the temperature was raised to room temperature, and subsequently the solution was stirred for 1 hour. After the completion of the reaction, the solution was concentrated under reduced pressure, and subsequently purified on silica gel column chromatography to yield 1.62 g as a colorless crystal having the following physical property.

Melting point: 104 to 107°C.

Example 13: Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-1-pyrazolylamide (Compound Number III-192)



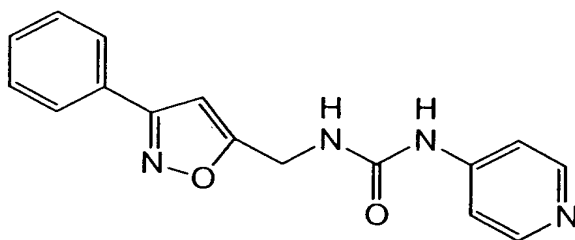
To a solution of 0.78 g of pyrazole in 30 ml of tetrahydrofuran, 0.93 of N,N'-carbonyl diimidazole was added

at room temperature, and the whole was stirred for one hour. Subsequently, 1.00 g of 5-aminomethyl-3-phenylisoxazole was added to the reaction solution, and the whole was stirred overnight. After the completion of the reaction, the reaction solution was concentrated under reduced pressure, and purified on silica gel column chromatography to yield 1.51 g as a colorless crystal having the following physical property.

Melting point: 137 to 138°C.

10

Example 14: Preparation of 3-[(3-phenyl-5-isoxazolyl)methyl]-1-(4-pyridyl) urea (Compound Number II-101)

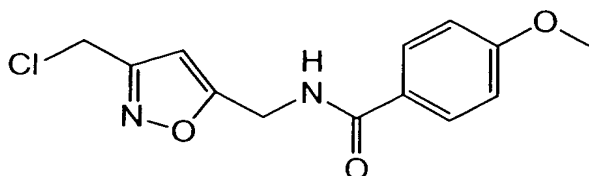


15 To a solution of 0.65 g of 4-aminopyridine in 30 ml of tetrahydrofuran, 0.93 of N,N'-carbonyl diimidazole was added at room temperature, and the whole was stirred for one hour. Subsequently, 1.00 g of 5-aminomethyl-3-phenylisoxazole was added to the reaction solution, and the whole was stirred  
20 overnight. After the completion of the reaction, the reaction solution was concentrated under reduced pressure, and purified on silica gel column chromatography to yield

1.62 g as a colorless crystal having the following physical property.

Melting point: 156 to 162°C.

- 5 Example 15: Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-methoxy benzamide (Compound Number III-3)

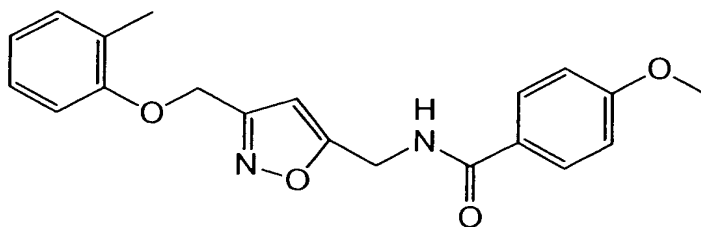


To a solution of 47.7 g of chloromethylaldoxime and  
10 6.43 g of N-(2-propynyl)-1-p-methoxy benzamide in 150 ml of dichloromethane, 380 g of 10% sodium hypochlorite aqueous solution was dripped over 1 hour at 0°C. The temperature was raised to room temperature, and then the solution was stirred overnight. Water was added, the solution was  
15 extracted twice with ethyl acetate, and an extract was washed with a saturated aqueous solution of sodium sulfite. The extract was dried on anhydrous magnesium sulfate, and subsequently purified on column chromatography to yield 5.83 g as a colorless solid having the following physical  
20 property.

Melting point: 93 to 110°C.

NMR :  $\delta$  3.86(3H,s), 4.53(2H,s), 4.72(2H,d), 6.31(1H,s), 6.69(1H,brs), 6.92(2H,d), 7.75(2H,d)

Example 16: Preparation of N-([3-(2-methylphenyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxy benzamide (Compound Number IV-47)



5 To a solution of 1.20 g of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-methoxy benzamide and 1.85 g of 2-methylphenol in 30 ml of dimethylformamide, 0.89 g of potassium carbonate was added at room temperature. The temperature was raised to 100°C, and the solution was  
10 stirred for 5 hours. Water was added, the solution was extracted twice with ethyl acetate, and a collected organic layer was washed with water and saturated sodium chloride solution. The organic layer was dried on anhydrous magnesium sulfate, and then purified on column  
15 chromatography to yield 0.65 g as a colorless solid having the following physical property.

Melting point: 115 to 117°C

NMR :  $\delta$  2.20(3H,s), 3.80(3H,s), 4.68(2H,d), 5.08(2H,s), 6.30(1H,s), 6.61-7.82(9H,m)

## 20 Example 17

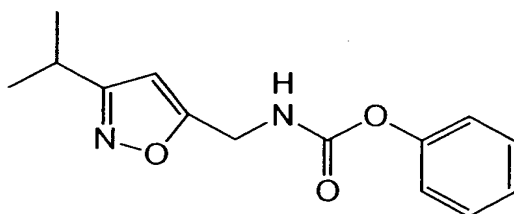
According to the methods described in Examples 1 to 16, the compounds in the following Examples 17(1) to 17(115)



were obtained.

Example 17(1): Production of [(3-isopropyl-5-isoxazolyl)methyl]phenylcarbamate (Compound number I-4):

5

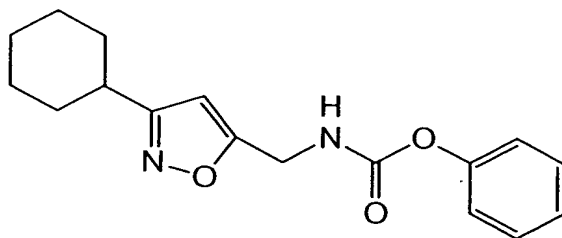


Property: oily;

NMR :  $\delta$  1.28(6H, d), 2.75-3.29(1H, m), 4.52(2H, d), 5.34-5.72(1H, brs), 6.12(1H, s), 7.00-7.48(5H, m)

Example 17(2): Preparation of phenyl [(3-cyclohexyl-5-isoxazolyl)methyl] carbamate (Compound number I-6):

10

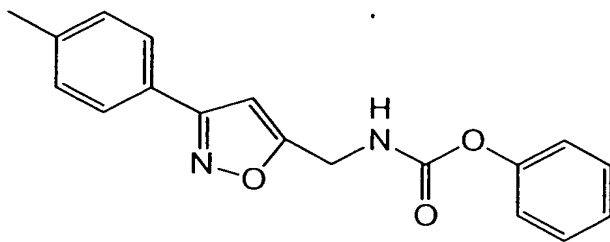


Property: paste;

NMR :  $\delta$  1.08-2.10(10H, m), 2.48-2.91(1H, brs), 4.42(2H, d), 5.80-6.00(1H, brs), 6.02(1H, s), 6.98-7.48(5H, m)

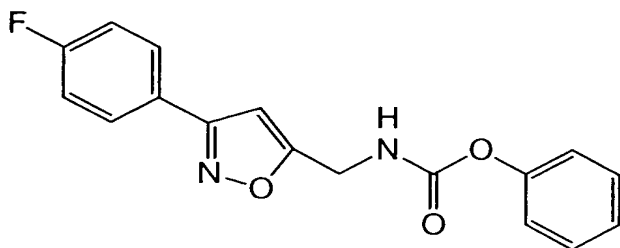
Example 17(3): Preparation of phenyl {[3-(4-tolyl)-5-isoxazolyl)methyl] carbamate (Compound number I-25):

15



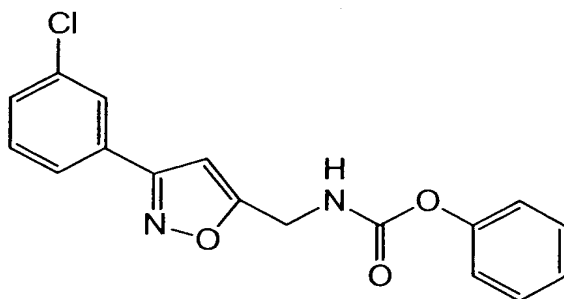
Melting point: 119 to 123°C.

Example 17(4): Preparation of phenyl {[3-(4-fluorophenyl)-  
5 5-isoxazolyl)methyl] carbamate (Compound number I-27):



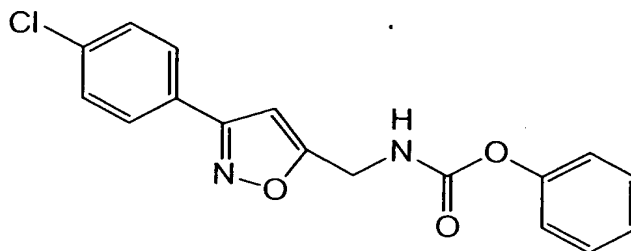
Melting point: 114 to 116°C.

Example 17(5): Preparation of phenyl {[3-(3-chlorophenyl)-  
10 5-isoxazolyl)methyl] carbamate (Compound number I-29):



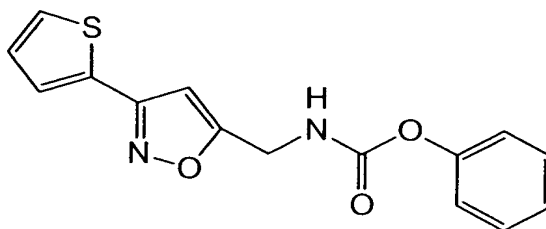
Melting point: 103 to 105°C.

Example 17(6): Preparation of phenyl {[3-(4-chlorophenyl)-  
15 5-isoxazolyl)methyl] carbamate (Compound number I-30):



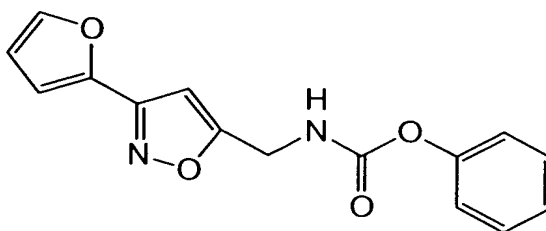
Melting point: 135 to 137°C.

Example 17(7): Preparation of phenyl {[3-(2-thienyl)-5-isoxazolyl)methyl] carbamate (Compound number I-32):



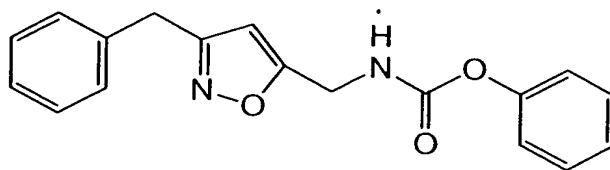
Melting point: 104 to 107°C.

Example 17(8): Preparation of phenyl {[3-(2-furyl)-5-isoxazolyl)methyl] carbamate (Compound number I-33):



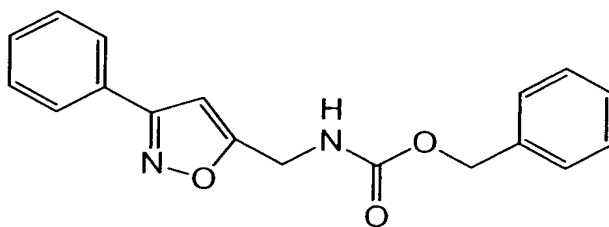
Melting point: 89 to 92°C.

Example 17(9): Preparation of phenyl [(3-benzyl-5-isoxazolyl)methyl] carbamate (Compound number I-38):



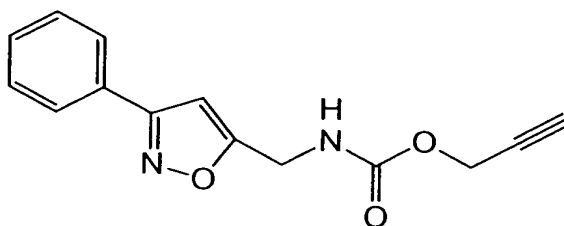
Melting point: 135 to 137°C.

Example 17(10): Preparation of benzyl [(3-phenyl-5-  
5 isoxazolyl)methyl] carbamate (Compound number I-41):



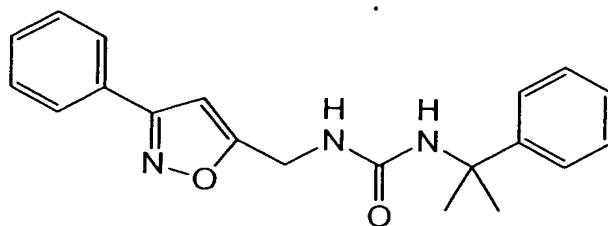
Melting point: 104 to 105°C.

Example 17(11): Preparation of [(3-phenyl-5-  
10 isoxazolyl)methyl]2-propynyl carbamate (Compound number I-42):



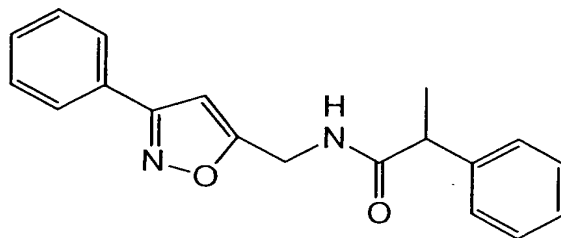
Melting point: 102 to 104°C.

15 Example 17(12): Preparation of 3-[(3-phenyl-5-  
isoxazolyl)methyl]-1-(methyl-1-phenylethyl)urea (Compound  
number I-45):



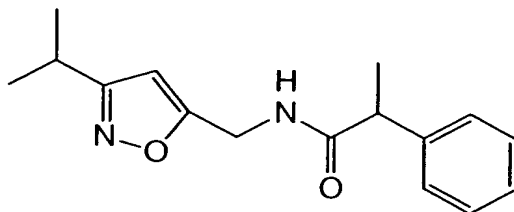
Melting point: 160 to 164°C.

Example 17(13): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2-phenylpropanamide (Compound number I-46):



Melting point: 110 to 112°C.

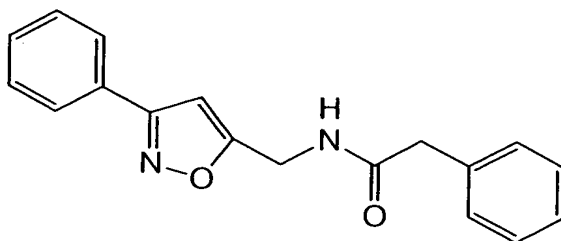
Example 17(14): Preparation of N-[(3-isopropyl-5-isoxazolyl)methyl]-2-phenylpropanamide (Compound number I-47):



Property: oily;

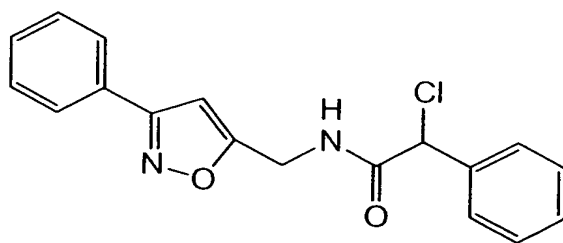
NMR :  $\delta$  0.80-1.66(9H, m), 2.67-3.14(1H, m), 3.27-3.81(1H, m), 4.40(2H, d), 5.80(1H, s), 6.22-6.51(1H, br), 7.26(5H, s)

Example 17(15): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2-phenylethanamide (Compound number I-48):



5 Melting point: 153 to 154°C.

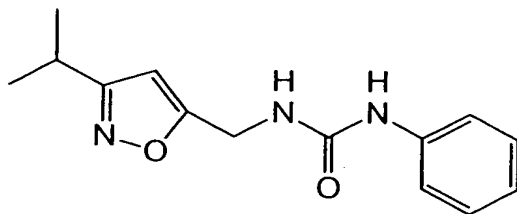
Example 17(16): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2-chloro-2-phenylethanamide (Compound number I-49):



10

Melting point: 121 to 125°C.

Example 17(17): Preparation of 1-[(3-isopropyl-5-isoxazolyl)methyl]-3-phenylurea (Compound number II-3):

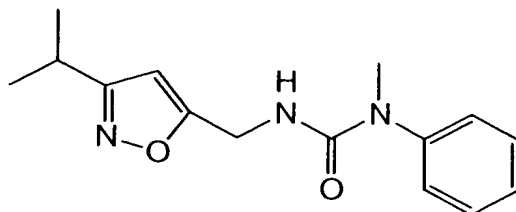


15

Property: paste;

NMR :  $\delta$  1.08(6H, d), 2.69-3.14(1H, m), 4.33(2H, d), 5.98(1H, s), 6.22-6.44(1H, brs), 6.86-7.31(5H, m), 7.92(1H, d)

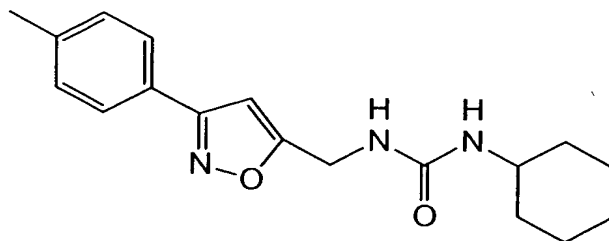
Example 17(18): Preparation of 1-[(3-isopropyl-5-isoxazolyl)methyl]-3-phenyl-3-methylurea (Compound number 5 II-4):



Property: oily;

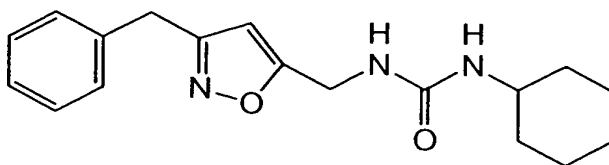
NMR :  $\delta$  1.24(6H, d), 2.72-3.21(1H, m), 4.42(2H, d), 4.77-5.04(1H, brs), 5.99(1H, s), 7.10-7.52(5H, m)

10 Example 17(19): Preparation of 1-[[3-(4-tolyl)-5-isoxazolyl]methyl]-3-cyclohexylurea (Compound number II-9):



Melting point: 163 to 170°C.

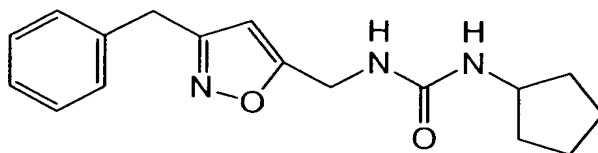
15 Example 17(20): Preparation of 1-[(3-benzyl-5-isoxazolyl)methyl]-3-cyclohexylurea (Compound number II-13):



Property: oily;

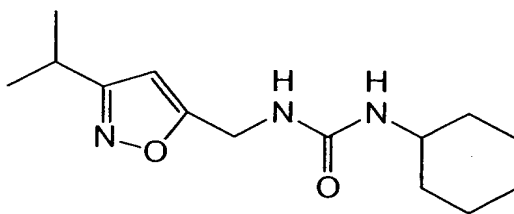
NMR :  $\delta$  0.80-2.05(10H, m), 3.22-3.68(1H, brs), 3.92(2H, s),  
4.35(2H, d), 5.23(1H, brt), 7.20(5H, brs)

Example 17(21): Preparation of 1-[(3-benzyl-5-  
5 isoxazolyl)methyl]-3-cyclopentylurea (Compound number II-  
14):



Melting point: 125 to 129°C.

10 Example 17(22): Preparation of 1-[(3-isopropyl-5-  
isoxazolyl)methyl]-3-cyclohexylurea (Compound number II-15):

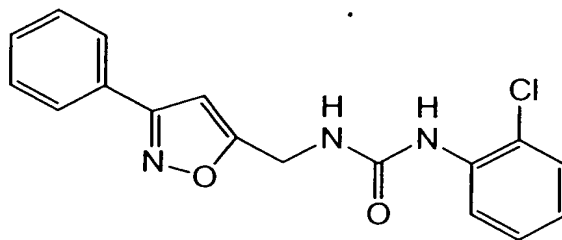


Property: oily;

15 NMR :  $\delta$  0.48-2.08(16H, m), 3.28-3.63(2H, m), 4.35-4.60(1H,  
brs), 4.43(2H, d), 4.90(1H, Brt), 6.12(1H, s)

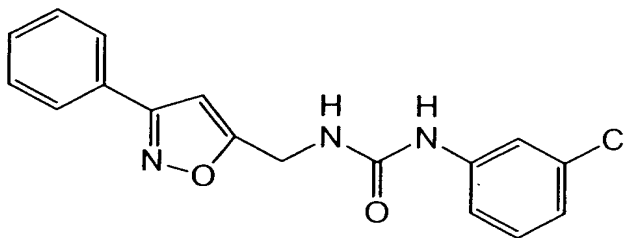
Example 17(23): Preparation of 1-[(3-phenyl-5-  
isoxazolyl)methyl]-3-(2-chlorophenyl)urea (Compound number  
II-48):





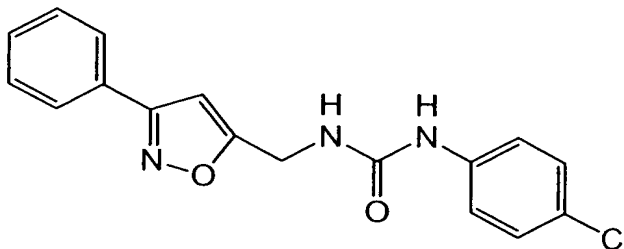
Melting point: 166 to 169°C.

Example 17(24): Preparation of 1-[(3-phenyl-5-isoxazolyl)methyl]-3-(3-chlorophenyl)urea (Compound number II-49):



NMR :  $\delta$  4.58 (2H, d) 6.51 (1H, s) 6.82-7.86 (10H, m) 8.08 (1H, brs).

10 Example 17(25): Preparation of 1-[(3-phenyl-5-isoxazolyl)methyl]-3-(4-chlorophenyl)urea (Compound number II-50):

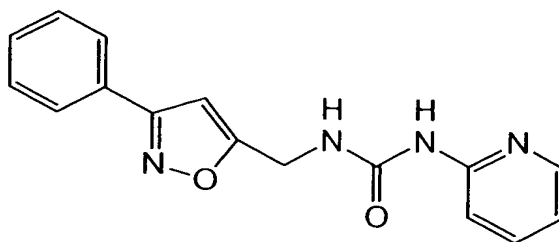


Melting point: 212 to 219°C.

15

Example 17(26): Preparation of 1-[(3-phenyl-5-

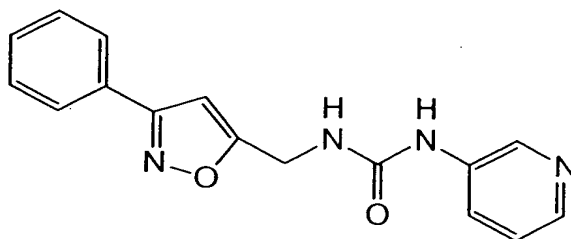
isoxazolyl)methyl]-3-(2-pyridyl)urea (Compound number II-99):



M.P.: 211 to 212°C.

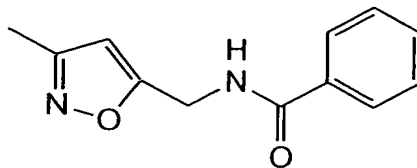
5

Example 17(27): Preparation of 1-[(3-phenyl-5-isoxazolyl)methyl]-3-(3-pyridyl)urea (Compound number II-100):



10 Melting point: 186 to 188°C.

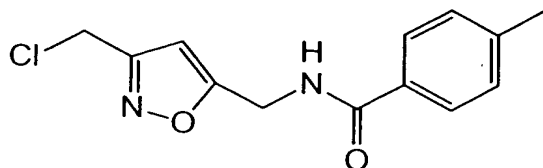
Example 17(28): Preparation of N-[(3-methyl-5-isoxazolyl)methyl]benzamide (Compound number III-1):



15 Melting point: 106 to 107°C.

Example 17(29): Preparation of N-[(3-chloromethyl-5-

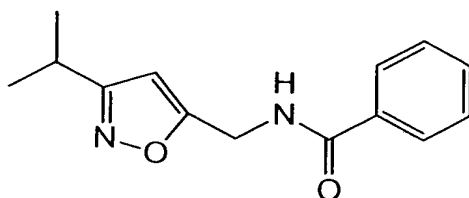
isoxazolyl)methyl]-4-methylbenzamide (Compound number III-2):



Property: paste;

5 NMR :  $\delta$  2.39(3H, s), 4.53(2H, s), 4.72(2H, d), 6.32(1H, s), 6.63-6.92(1H, brs), 7.21(2H, d), 7.68(2H, d)

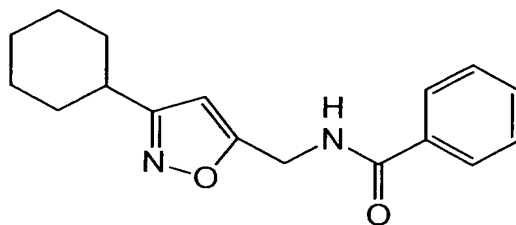
Example 17(30): Preparation of N-[(3-isopropyl-5-isoxazolyl)methyl]benzamide (Compound number III-4):



10 Property: oily;

NMR :  $\delta$  1.25(6H, d), 2.80-3.27(1H, m), 4.71(2H, d), 6.09(1H, s), 6.66(1H, brs), 7.28-7.88(5H, m)

Example 17(31): Preparation of N-[(3-cyclohexyl-5-isoxazolyl)methyl]benzamide (Compound number III-6):

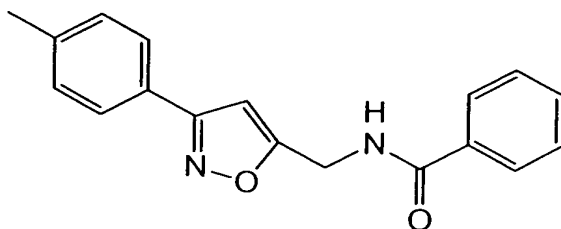


15

Melting point: 89 to 93°C.

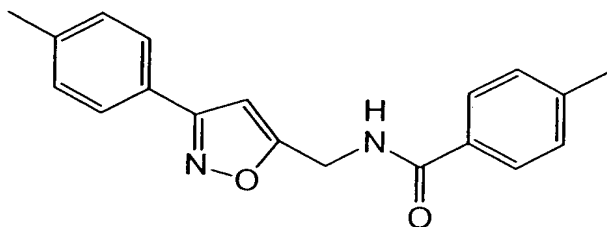
Example 17(32): Preparation of N-[(3-(4-tolyl)-5-

isoxazolyl)methyl}benzamide (Compound number III-10):



Melting point: 154 to 156°C.

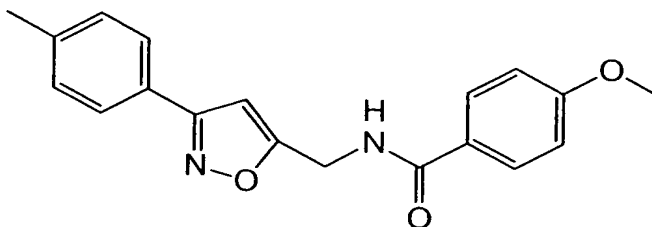
5 Example 17(33): Preparation of N-([3-(4-tolyl)-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number III-11):



Melting point: 164 to 166°C.

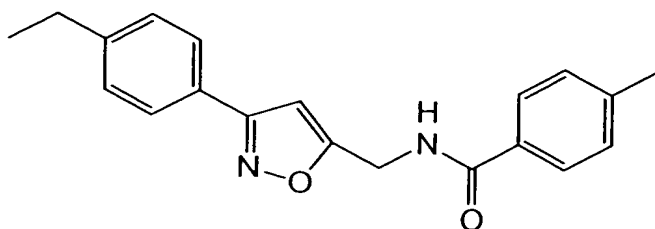
10

Example 17(34): Preparation of N-([3-(4-tolyl)-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-12):



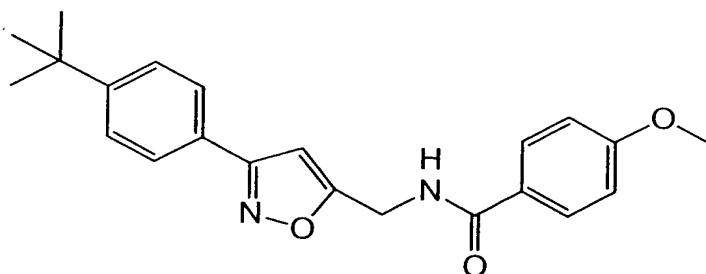
15 Melting point: 174 to 175°C.

Example 17(35): Preparation of N-[[3-(4-ethylphenyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number III-13):



5 Melting point: 150 to 152°C.

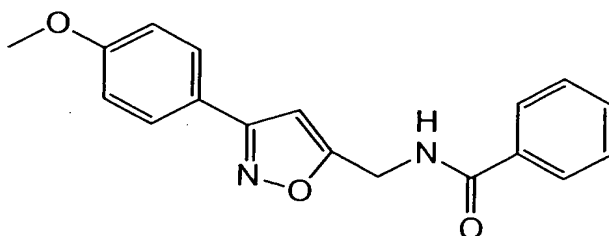
Example 17(36): Preparation of N-[[3-(4-t-butylphenyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number III-14):



10

Melting point: 179 to 180°C.

Example 17(37): Preparation of N-[[3-(4-methoxyphenyl)-5-isoxazolyl]methyl]benzamide (Compound number III-17):

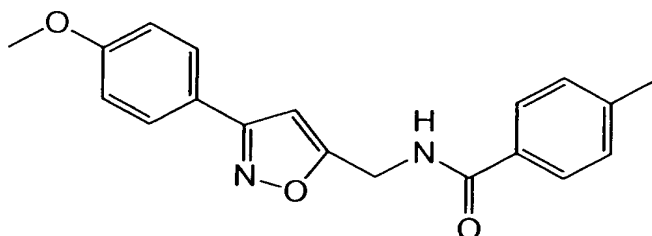


15

Melting point: 125 to 127°C.

Example 17(38): Preparation of N-[[3-(4-methoxyphenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-19):

5

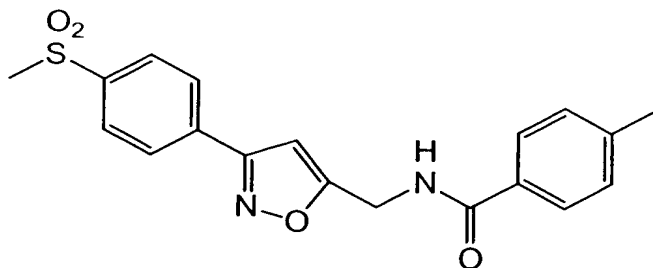


Melting point: 139 to 142°C;

NMR :  $\delta$  2.37(3H, s), 3.81(3H, s), 4.75(2H, d), 6.43(1H, s), 6.69(1H, brs), 6.82-7.83(8H, m)

Example 17(39): Preparation of N-[[3-(4-mesyphenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-26):

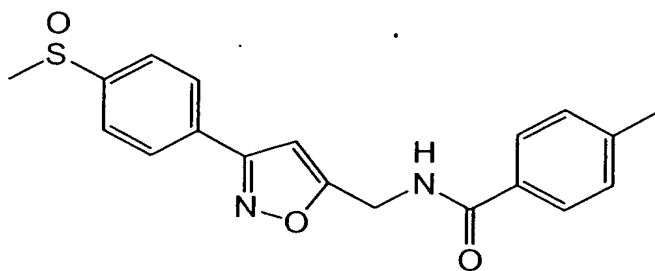
10



NMR :  $\delta$  2.37(3H, s), 3.08(3H, s), 4.78(2H, d), 6.61(1H, s), 7.09-8.08(9H, m)

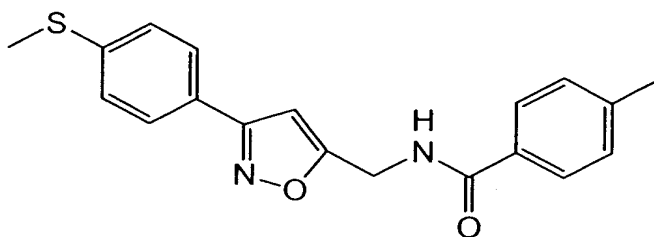
Example 17(40): Preparation of N-[[3-(4-methanesulfinyl phenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-27):

15



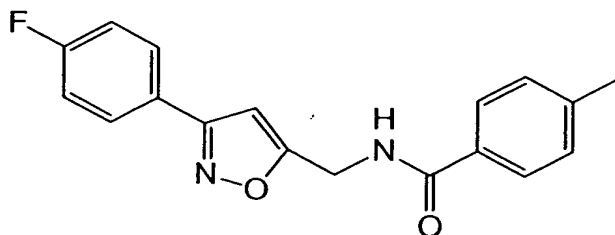
NMR :  $\delta$  2.37(3H, s), 2.72(3H, s), 4.78(2H, d), 5.99(1H, s), 7.09-8.08(9H, m)

Example 17(41): Preparation of N-[[3-(4-methylthiophenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-28):



Melting point: 135 to 155°C.

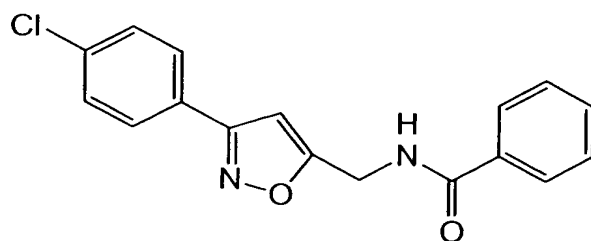
10 Example 17(42): Preparation of N-[[3-(4-fluorophenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-31):



Melting point: 167 to 168°C;

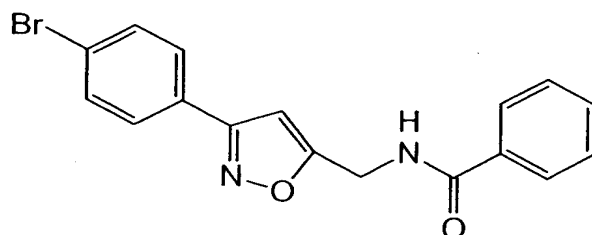
15 NMR :  $\delta$  2.67(3H, s), 4.78(2H, d), 6.50(1H, s), 6.60-7.93(9H, m)。

Example 17(43): Preparation of N-([3-(4-chlorophenyl)-5-isoxazolyl]methyl)benzamide (Compound number III-34):



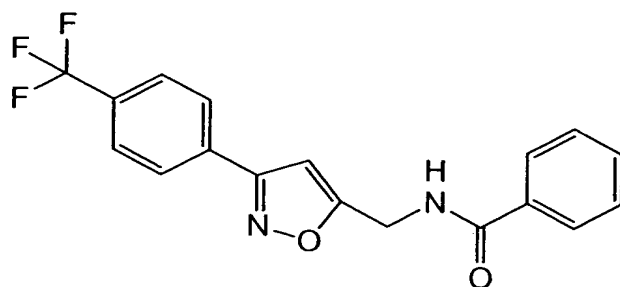
5 Melting point: 172 to 173°C.

Example 17(44): Preparation of N-([3-(4-bromophenyl)-5-isoxazolyl]methyl)benzamide (Compound number III-37):



10 Melting point: 184 to 187°C.

Example 17(45): Preparation of N-([3-(4-trifluoromethylphenyl)-5-isoxazolyl]methyl)benzamide (Compound number III-40):

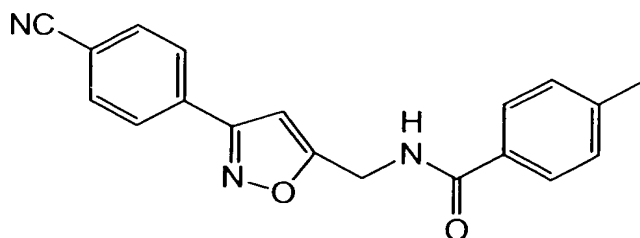


15



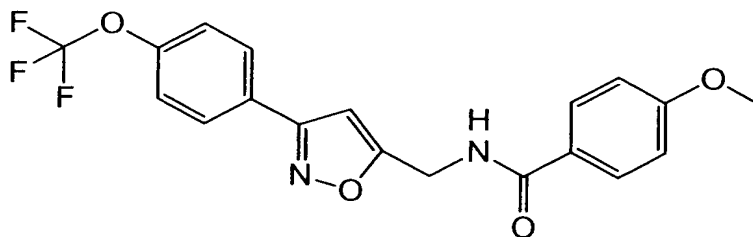
Melting point: 179 to 180°C.

Example 17(46): Preparation of N-([3-(4-cyanophenyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-43):



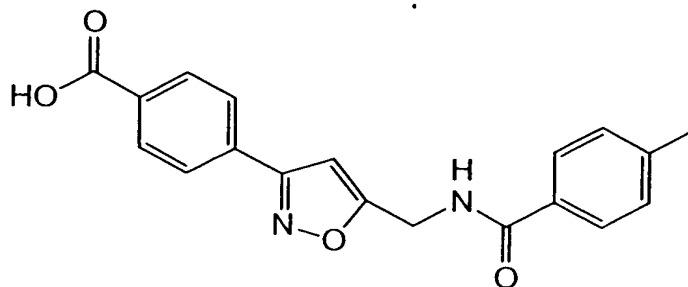
NMR :  $\delta$  2.33 (3H, s), 4.79 (2H, d), 6.44-6.85 (1H, br), 6.59 (1H, s), 7.10-8.87 (8H, m)

Example 17(47): Preparation of N-([3-(4-trifluoromethoxyphenyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-49):



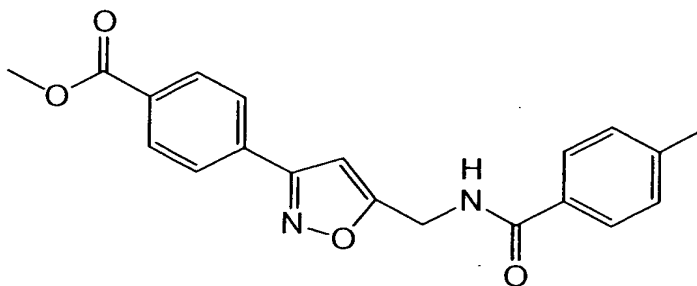
Melting point: 188 to 191°C.

Example 17(48): Preparation of N-([3-(4-carboxyphenyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-51):



Melting point: 248°C or more (degradation).

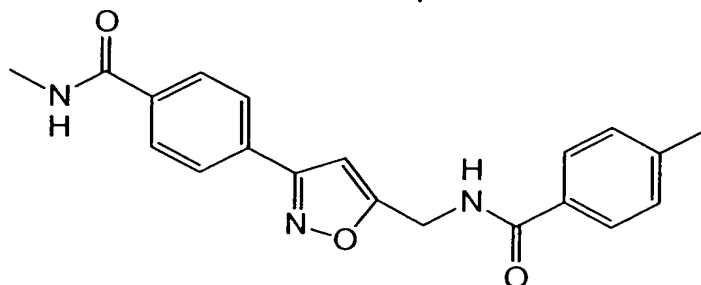
Example 17(49): Preparation of N-{{3-(4-methoxycarbonylphenyl)-5-isoxazolyl}methyl}-4-methylbenzamide (Compound number III-52):



Melting point: 173 to 175°C;

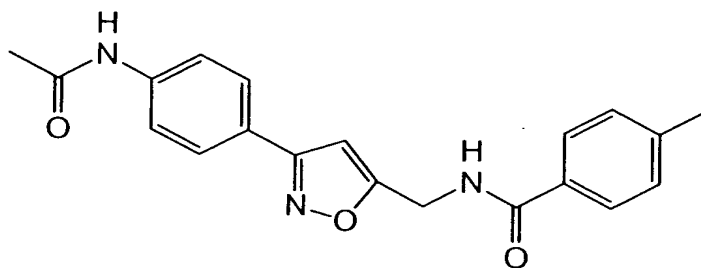
NMR :  $\delta$  2.34(3H, s), 3.92(3H, d), 4.76(2H, d), 6.53(1H, s),  
 10 6.92-8.15(9H, m).

Example 17(50): Preparation of N-{{3-(4-methylaminocarbonylphenyl)-5-isoxazolyl}methyl}-4-methylbenzamide (Compound number III-54):



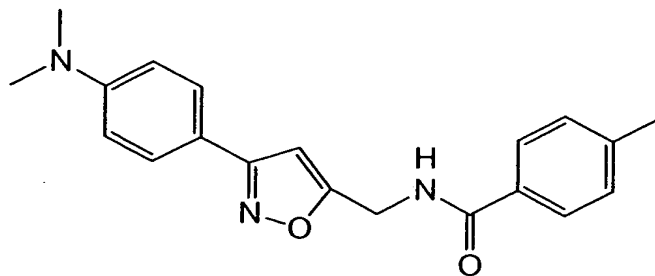
Melting point: 261 to 262°C.

Example 17(51): Preparation of N-[[3-(4-acetamidophenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-57):



Melting point: > 200°C.

10 Example 17(52): Preparation of N-[[3-(4-dimethylaminophenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-61):

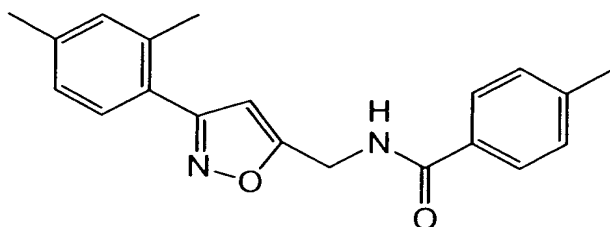


NMR :  $\delta$  2.33 (3H, s), 2.99 (6H, s), 4.74 (2H, d), 6.46 (1H, s),

6.53-8.07 (9H, m)

Example 17(53): Preparation of N-[[3-(2,4-dimethylphenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-65):

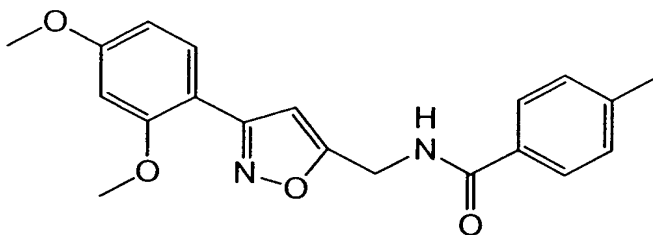
5



Melting point: 138 to 140°C.

Example 17(54): Preparation of N-[[3-(2,4-dimethoxyphenyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number III-66):

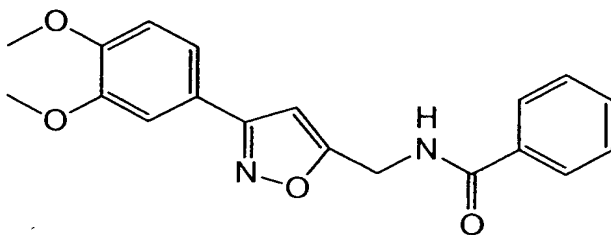
10



Melting point: 157 to 158°C.

Example 17(55): Preparation of N-[[3-(3,4-dimethoxyphenyl)-5-isoxazolyl]methyl]benzamide (Compound number III-67):

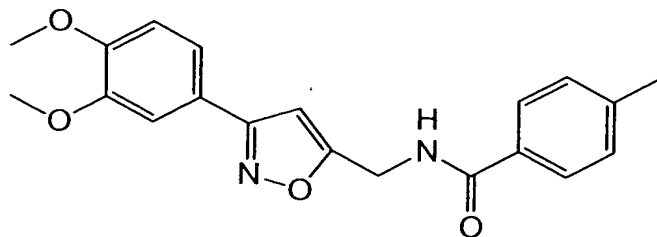
15



Melting point: 122 to 130°C.

Example 17(56): Preparation of N-{{[3-(3,4-dimethoxyphenyl)-5-isoxazolyl]methyl}-4-methylbenzamide (Compound number III-

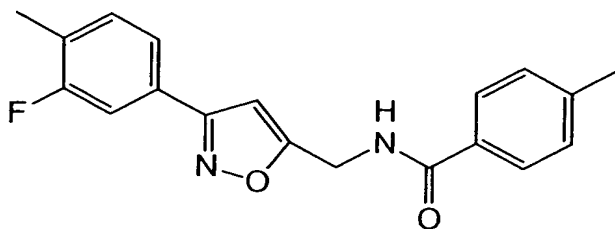
5 68):



Melting point: 160 to 162°C;

NMR :  $\delta$  2.32(3H, s), 3.82(6H, s), 4.72(2H, d), 6.43(1H, s), 6.72-7.83(8H, m)

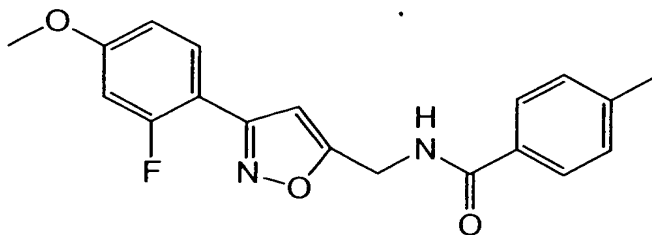
10 Example 17(57): Preparation of N-{{[3-(3-fluoro-4-methylphenyl)-5-isoxazolyl]methyl}-4-methylbenzamide (Compound number III-73):



Melting point: 165 to 166°C.

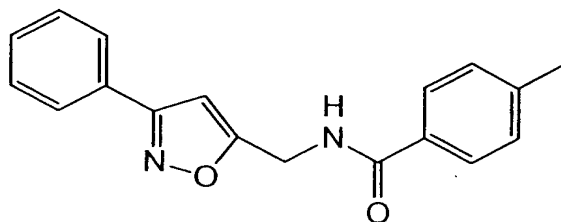
15

Example 17(58): Preparation of N-{{[3-(2-fluoro-4-methoxyphenyl)-5-isoxazolyl]methyl}-4-methylbenzamide (Compound number III-74):



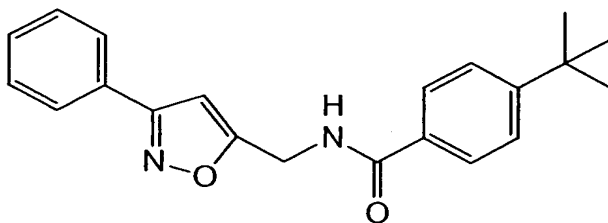
Melting point: 154 to 155°C.

Example 17(59): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number III-107):

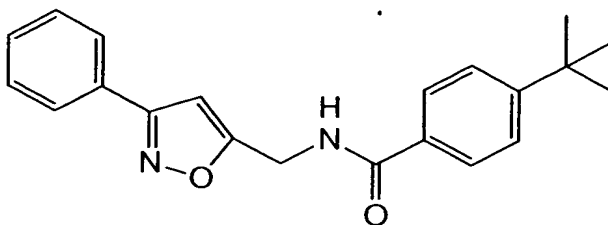


Melting point: 154 to 156°C.

10 Example 17(60): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-t-butylbenzamide (Compound number III-108):

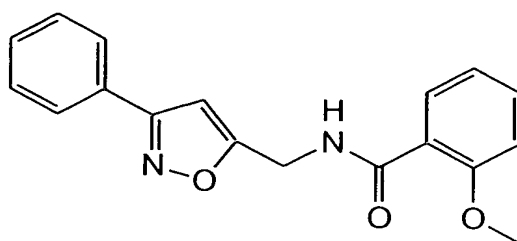


Melting point: 135 to 140°C.



Example 17(61): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2-methoxybenzamide (Compound number III-109):

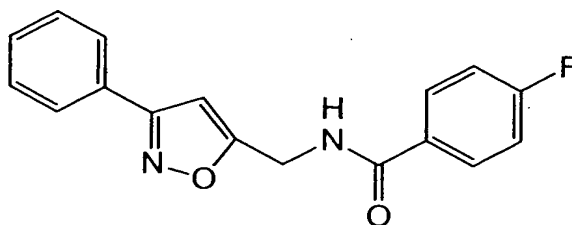
5



Property: oily;

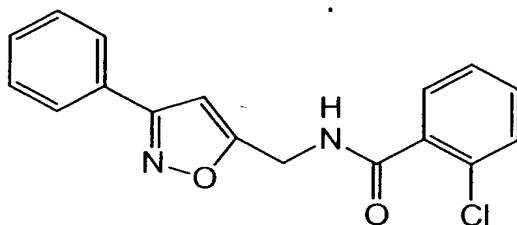
NMR :  $\delta$  4.00(3H, s) 4.81(2H, d) 6.53(1H, s) 6.87-8.55(10H, m)

Example 17(62): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-fluorobenzamide (Compound number III-10 121):



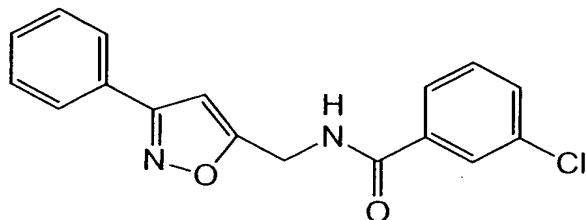
Melting point: 126 to 131°C.

Example 17(63): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2-chlorobenzamide (Compound number III-15 122):



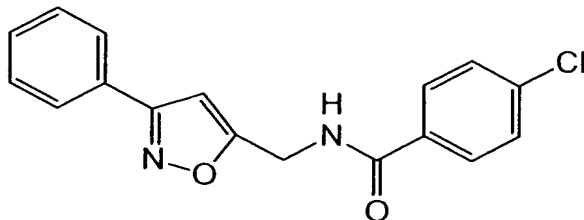
Melting point: 170 to 175°C.

Example 17(64): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-3-chlorobenzamide (Compound number III-123):



Melting point: 113 to 115°C.

Example 17(65): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-124):

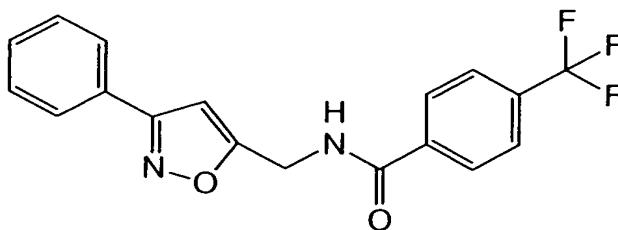


Melting point: 148 to 151°C.

Example 17(66): Preparation of N-[(3-phenyl-5-



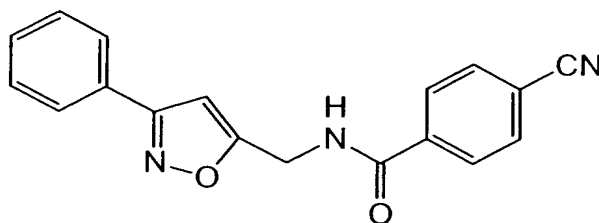
isoxazolyl)methyl]-4-trifluoromethylbenzamide (Compound number III-130):



Melting point: 157 to 168°C.

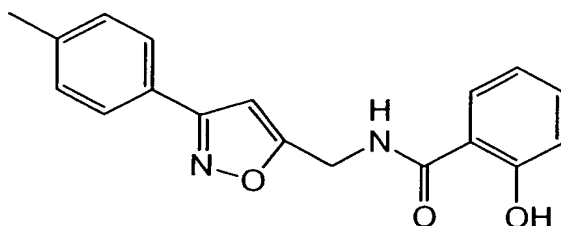
5

Example 17(67): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-cyanobenzamide (Compound number III-133):



10 Melting point: 144 to 145°C.

Example 17(68): Preparation of N-[(3-(4-tolyl)-5-isoxazolyl)methyl]-2-hydroxybenzamide (Compound number III-134):

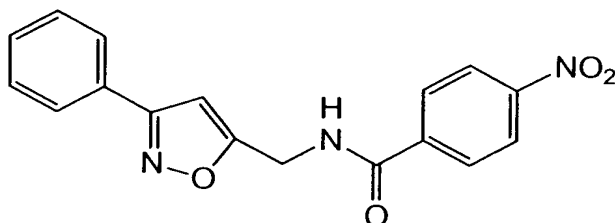


15

Melting point: 120 to 132°C.

Example 17(69): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-nitrobenzamide (Compound number III-137):

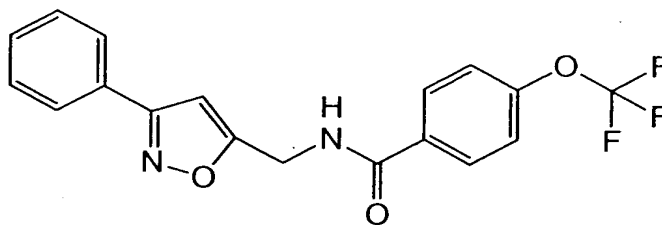
5



Melting point: 166 to 174°C.

Example 17(70): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-4-trifluoromethoxybenzamide (Compound number III-139):

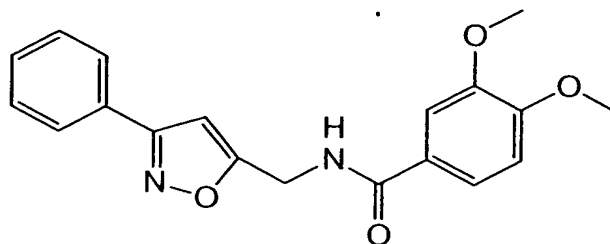
10



Melting point: 140 to 141°C.

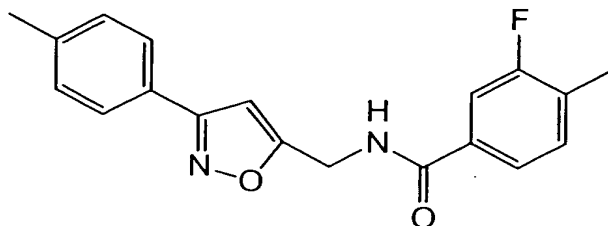
Example 17(71): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-3,4-dimethoxybenzamide (Compound number III-156):

15



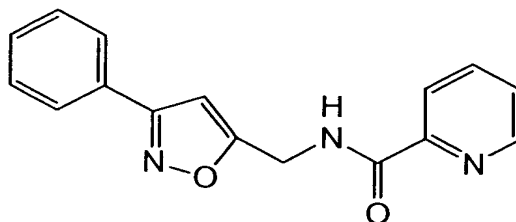
Melting point: 157 to 169°C.

Example 17(72): Preparation of N-[(3-(4-tolyl)-5-isoxazolyl)methyl]-3-fluoro-4-methylbenzamide (Compound number III-162):



Melting point: 148 to 154°C.

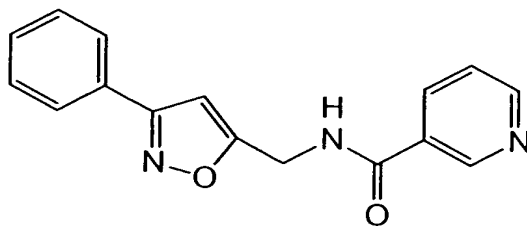
Example 17(73): Preparation of (3-phenyl-5-isoxazolyl)methylaminocarbonyl-2-pyridine (Compound number III-174):



Melting point: 125 to 127°C.

Example 17(74): Preparation of (3-phenyl-5-

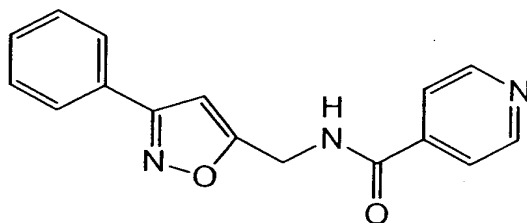
isoxazolyl)methylaminocarbonyl-3-pyridine (Compound number III-175):



Melting point: 130 to 132°C.

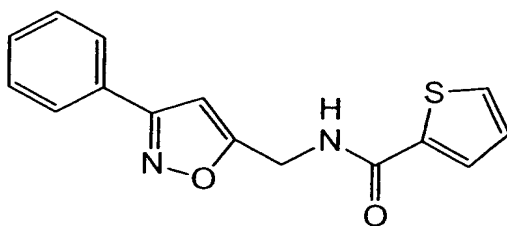
5

Example 17(75): Preparation of (3-phenyl-5-isoxazolyl)methylaminocarbonyl-4-pyridine (Compound number III-176):



10 Melting point: 125 to 135°C.

Example 17(76): Preparation of (3-phenyl-5-isoxazolyl)methylaminocarbonyl-2-thiophene (Compound number III-177):

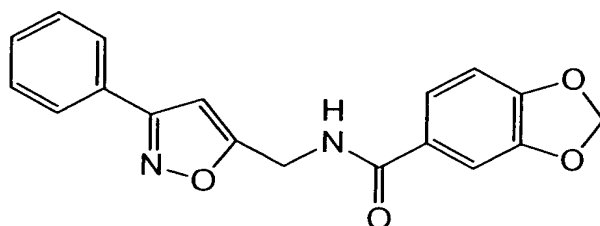


15

Melting point: 141 to 155°C.

Example 17(77): Preparation of 5-(3-phenyl-5-isoxazolyl)methylaminocarbonyl-1,3-benzodioxol (Compound number III-180):

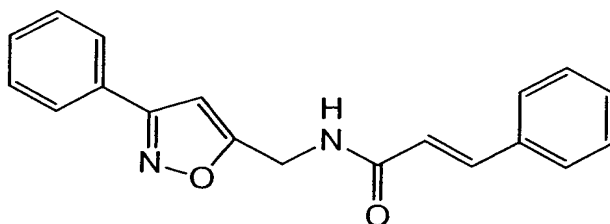
5



Melting point: 175 to 183°C.

Example 17(78): Preparation of (E)-N-[(3-phenyl-5-isoxazolyl)methyl]-3-phenylpropenamide (Compound number III-201):

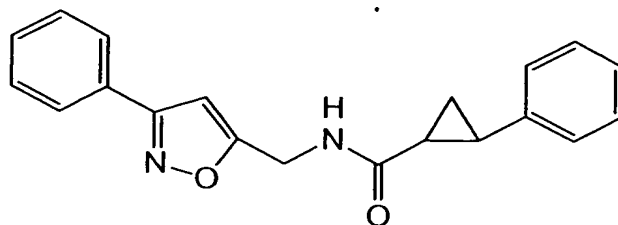
10



Melting point: 155 to 157°C.

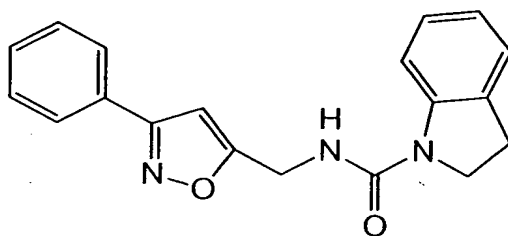
Example 17(79): Preparation of 1-[(3-phenyl-5-isoxazolyl)methylaminocarbonyl]-2-phenylcyclopropane (Compound number III-202):

15



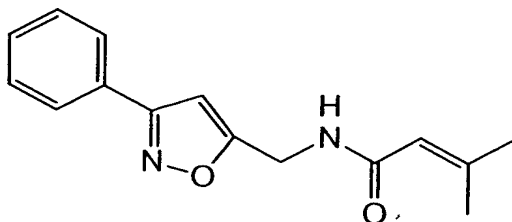
Melting point: 152 to 154°C.

Example 17(80): Preparation of N-[(3-phenyl-5-isoxazolyl)methylaminocarbonyl]indoline (Compound number III-208):



Melting point: 137 to 139°C.

10 Example 17(81): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-3-methyl-2-butenamide (Compound number III-209):

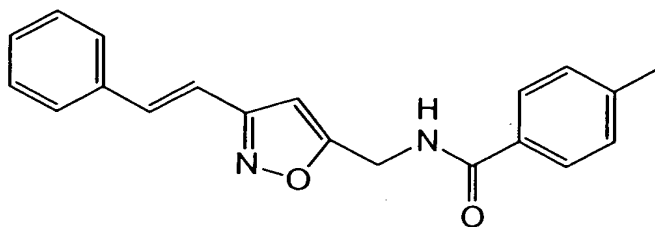


Melting point: 97 to 98°C.

15

Example 17(82): Preparation of N-[[3-(2-phenylethenyl)-5-

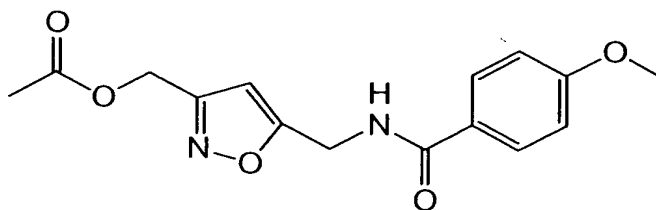
isoxazolyl)methyl]-4-methylbenzamide (Compound number III-210):



Melting point: 151 to 156°C.

5

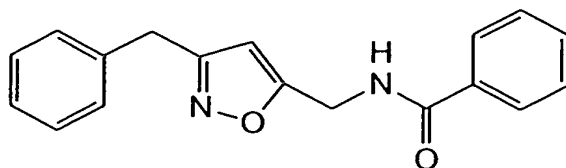
Example 17(83): Preparation of N-[(3-acetyloxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-211):



10 Property: paste;

NMR :  $\delta$  2.09(3H, s), 3.80(3H, s), 4.68(2H, d), 5.09(2H, d), 6.21(1H, s), 6.87(2H, d), 7.19-7.40(1H, brt), 7.78(2H, d)

Example 17(84): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]benzamide (Compound number IV-1):



15

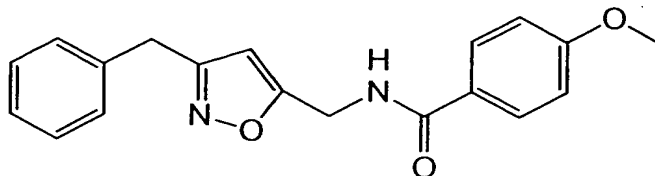
Property: oily;

NMR :  $\delta$  3.92(2H, s), 4.61(2H, d), 5.94(1H, s), 6.83-7.88(11H,

m)

Example 17(85): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-2):

5

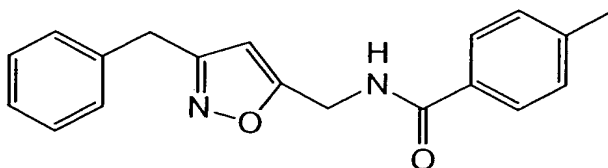


Property: oily;

NMR :  $\delta$  3.80(3H, s), 3.93(2H, s), 4.61(2H, d), 5.98(1H, s), 6.86(2H, d), 7.21(5H, s), 7.73(2H, d)

Example 17(86): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number IV-3):

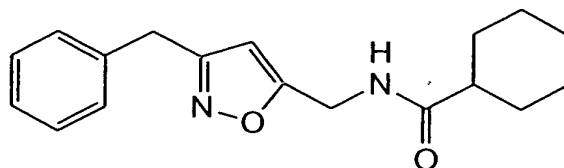
10



Melting point: 119 to 124°C.

Example 17(87): Preparation of (3-benzyl-5-isoxazolyl)methylaminocarbonylcyclohexane (Compound number IV-4):

15



Property: oily;

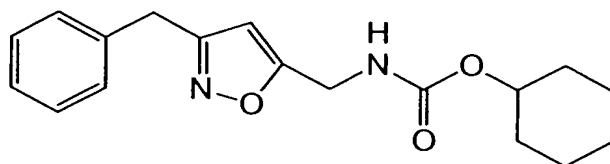
NMR :  $\delta$  1.06-2.30(11H, m), 3.99(2H, s), 4.50(2H, d), 5.71-



6.04(1H, br), 5.89(1H, s), 7.21(5H, s)

Example 17(88): Preparation of [(3-benzyl-5-isoxazolyl)methyl]cyclohexylcarbamate (Compound number IV-5):

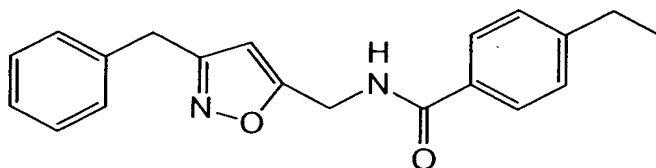
5



Melting point: 98 to 104°C.

Example 17(89): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-4-ethylbenzamide (Compound number IV-6):

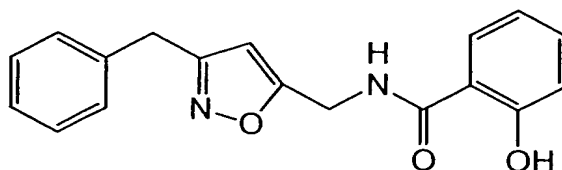
10



Property: oily;

NMR :  $\delta$  1.22(3H, t), 2.70(2H, q), 3.93(2H, s), 4.67(2H, d), 5.99(1H, s), 6.40-6.72(1H, brs), 7.08-7.47(7H, m), 7.69(2H, d)

15 Example 17(90): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-2-hydroxybenzamide (Compound number IV-7):

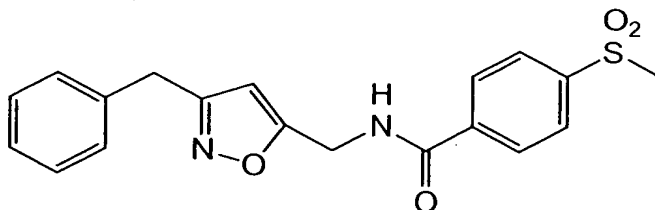


Property: paste;

NMR :  $\delta$  3.92 (2H, s), 4.60 (2H, d), 5.99 (1H, s), 6.47-7.56 (10H, m), 11.4 (1H, s)

Example 17(91): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-4-methanesulfonylbenzamide (Compound

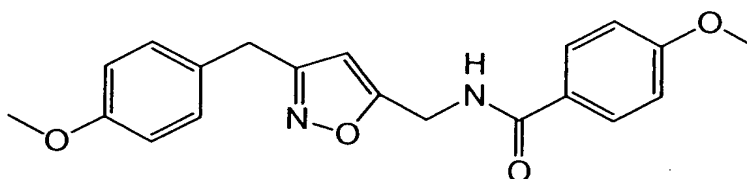
5 number IV-8):



Melting point: 148 to 157°C;

NMR :  $\delta$  3.04 (3H, s), 3.97 (2H, s), 4.69 (2H, d), 6.01 (1H, s), 6.95-7.55 (6H, m), 7.93 (4H, s)

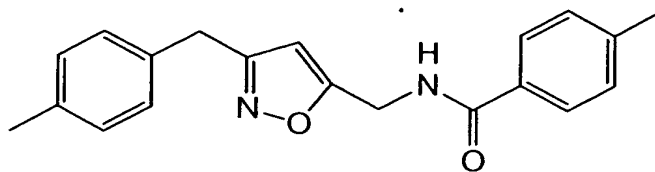
10 Example 17(92): Preparation of N-[[3-(4-methoxybenzyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-9):



Melting point: 114 to 117°C.

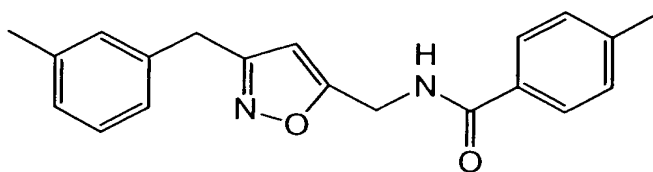
15

Example 17(93): Preparation of N-[[3-(4-methylbenzyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number IV-10):



Melting point: 86 to 97°C.

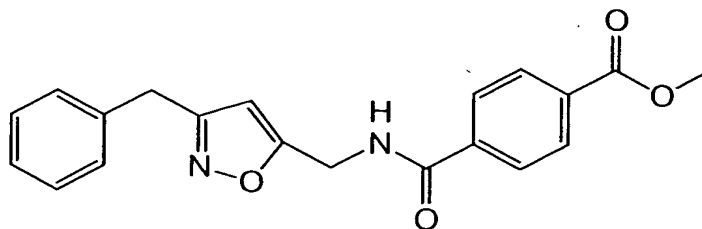
Example 17(94): Preparation of N-[[3-(3-methylbenzyl)-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number IV-11):



Property: oily;

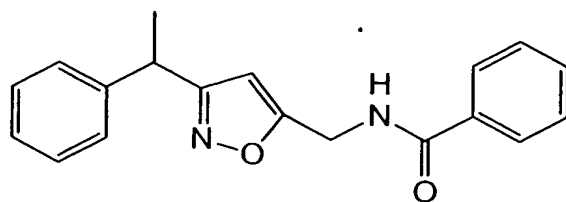
NMR :  $\delta$  2.31(3H, s), 2.40(3H, s), 3.92(2H, s), 4.65(2H, d), 5.99(1H, s), 6.53-6.87(1H, brs), 6.87-7.86(8H, m)

Example 17(95): Preparation of N-[(3-benzyl-5-isoxazolyl)methyl]-4-methoxycarbonylbenzamide (Compound number IV-23):



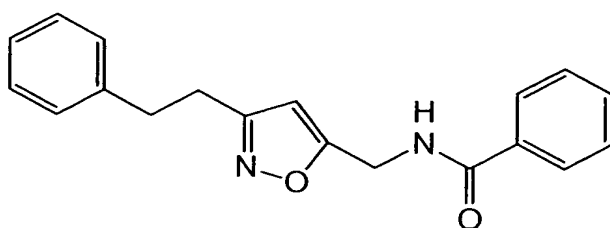
Melting point: 134 to 140°C.

Example 17(96): Preparation of N-[[3-(1-phenylethyl)-5-isoxazolyl]methyl]benzamide (Compound number IV-29):



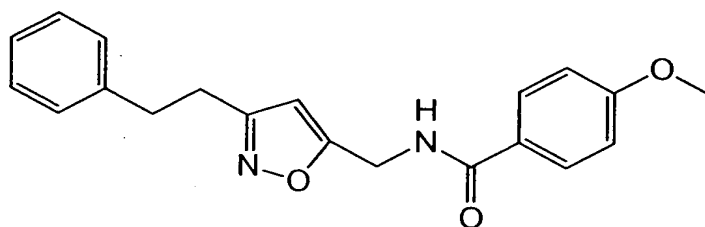
Melting point: 100 to 107°C.

Example 17(97): Preparation of N-([3-(2-phenylethyl)-5-  
5 isoxazolyl]methyl)benzamide (Compound number IV-33):



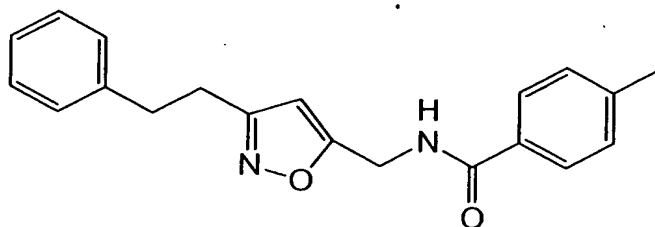
Melting point: 53 to 83°C.

Example 17(98): Preparation of N-([3-(2-phenylethyl)-5-  
10 isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-34):



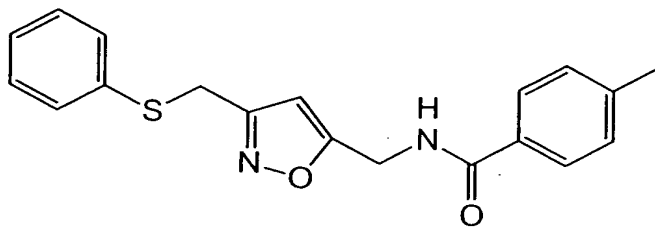
Melting point: 119 to 131°C.

15 Example 17(99): Preparation of N-([3-(2-phenylethyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number IV-35):



Melting point: 131 to 137°C.

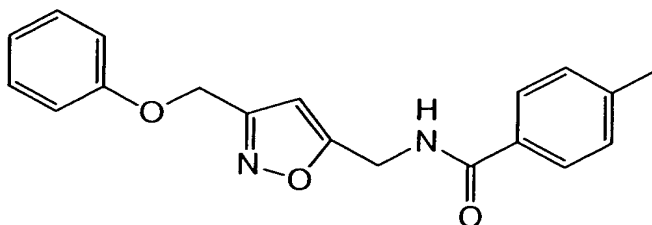
Example 17(100): Preparation of N-[(3-phenylthiomethyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number IV-40):



Melting point: 79 to 85°C;

10 NMR :  $\delta$  2.40(3H, s), 4.04(2H, s), 4.62(2H, d), 6.13(1H, s), 6.76(1H, s), 7.03-7.74(10H, m)

Example 17(101): Preparation of N-[(3-phenoxyethyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number IV-46):



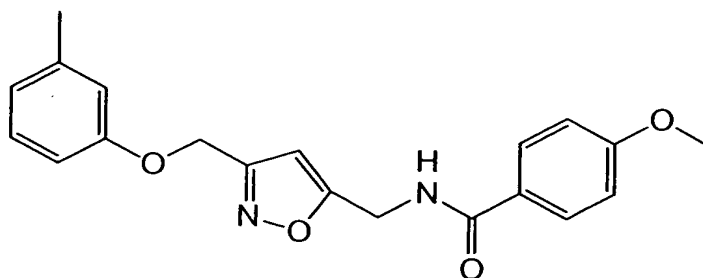
15 Property: paste;

NMR :  $\delta$  2.36(3H, s), 4.67(2H, d), 5.07(2H, s), 6.30(1H, s),

6.52-7.78 (10H, m)

Example 17(102): Preparation of N-[[3-(3-methylphenyl)oxymethyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-48):

5

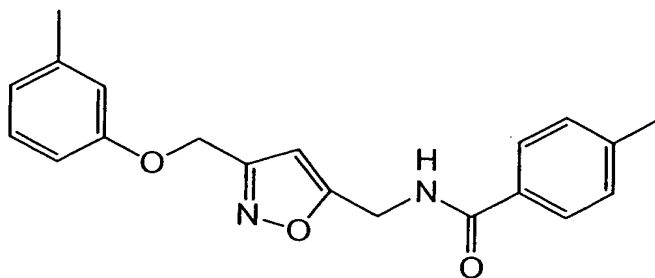


Melting point: 86 to 90°C;

NMR :  $\delta$  2.25 (3H, s), 3.75 (3H, s), 4.62 (2H, s), 5.00 (2H, s), 6.27 (1H, s), 6.60-7.83 (9H, m)

Example 17(103): Preparation of N-[[3-(3-methylphenyl)oxymethyl-5-isoxazolyl]methyl]-4-methylbenzamide (Compound number IV-49):

10

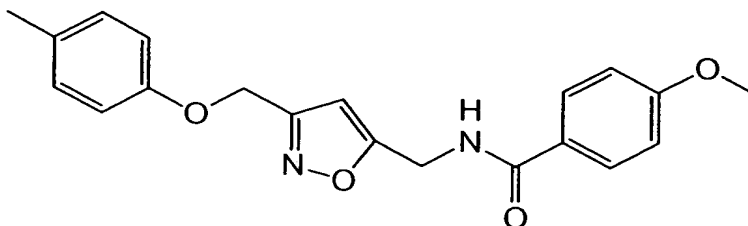


Melting point: 94 to 96°C;

15 NMR :  $\delta$  2.31 (3H, s), 2.35 (3H, s), 4.72 (2H, d), 5.05 (2H, s), 6.33 (1H, s), 6.61-7.76 (9H, m)

Example 17(104): Preparation of N-[[3-(4-methylphenyl)oxymethyl-5-isoxazolyl]methyl]-4-

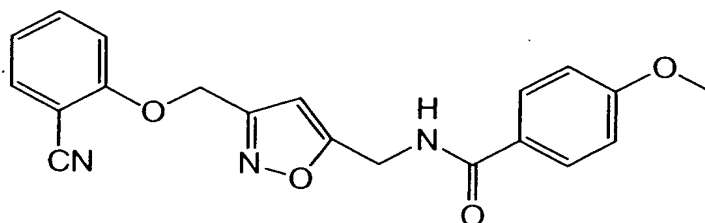
methoxybenzamide (Compound number IV-50):



Melting point: 124 to 127°C;

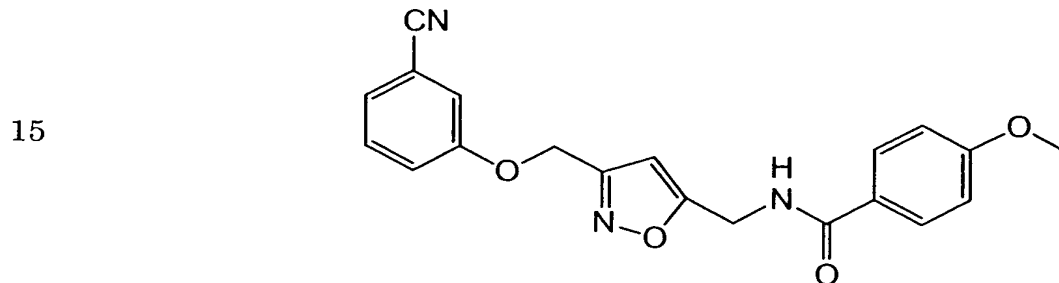
5 NMR :  $\delta$  2.27(3H, s), 3.80(3H, s), 4.68(2H, s), 5.03(2H, s),  
6.33(1H, s), 6.64-8.00(9H, m)

Example 17(105): Preparation of N-([3-(2-cyanophenyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-51):



10 Melting point: 111 to 114°C.

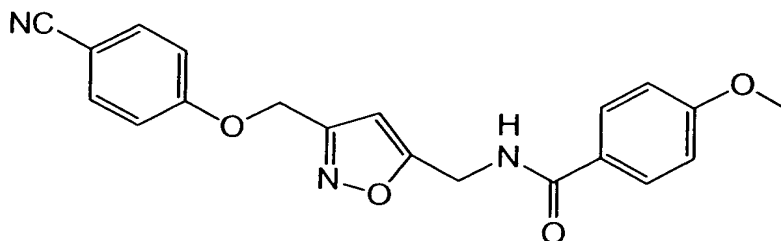
Example 17(106): Preparation of N-([3-(3-cyanophenyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-52):



Melting point: 128 to 130°C.

Example 17(107): Preparation of N-{{3-(4-cyanophenyl)oxymethyl-5-isoxazolyl)methyl}-4-

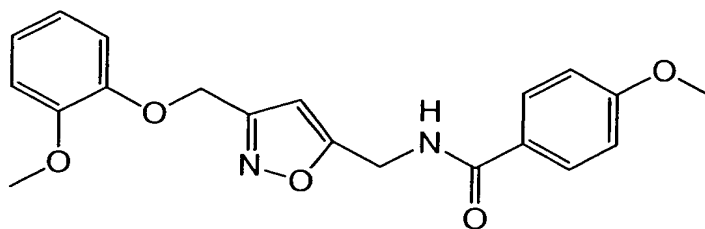
5 methoxybenzamide (Compound number IV-53):



Melting point: 119 to 121°C;

NMR :  $\delta$  3.81(3H, s), 4.72(2H, d), 5.14(2H, s), 6.32(1H, s), 6.73-8.02(9H, m)

10 Example 17(108): Preparation of N-{{3-(2-methoxyphenyl)oxymethyl-5-isoxazolyl)methyl}-4-methoxybenzamide (Compound number IV-54):

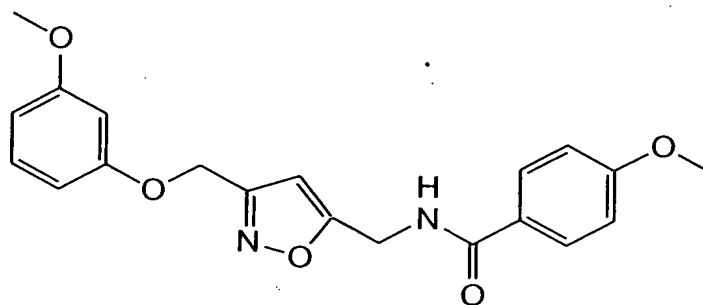


Property: oily;

15 NMR :  $\delta$  3.75(6H, s), 4.60(2H, d), 5.07(2H, s), 6.28(1H, s), 6.69-7.90(9H, m)

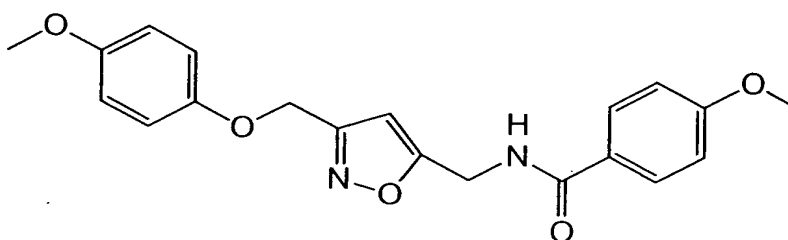
Example 17(109): Preparation of N-{{3-(3-methoxyphenyl)oxymethyl-5-isoxazolyl)methyl}-4-methoxybenzamide (Compound number IV-55):





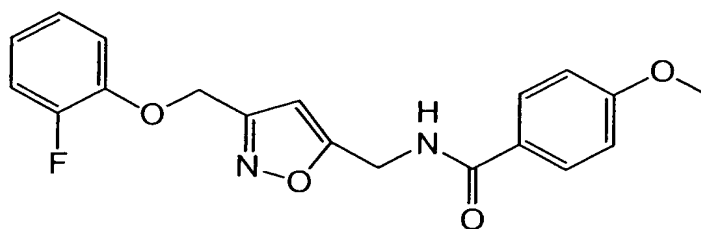
Melting point: 95 to 104°C.

Example 17(110): Preparation of N-([3-(4-methoxyphenyl)oxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-56):



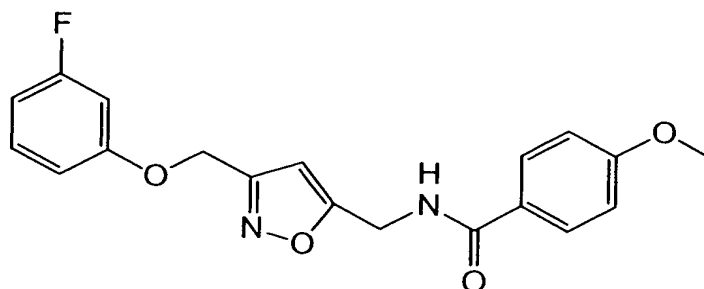
Melting point: 115 to 121°C.

10 Example 17(111): Preparation of N-([3-(2-fluorophenyl)oxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-57):



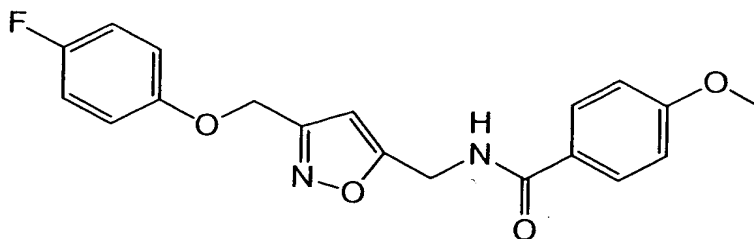
Melting point: 97 to 102°C.

Example 17(112): Preparation of N-{{3-(3-fluorophenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-58):



5 Melting point: 95 to 106°C.

Example 17(113): Preparation of N-{{3-(4-fluorophenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-59):

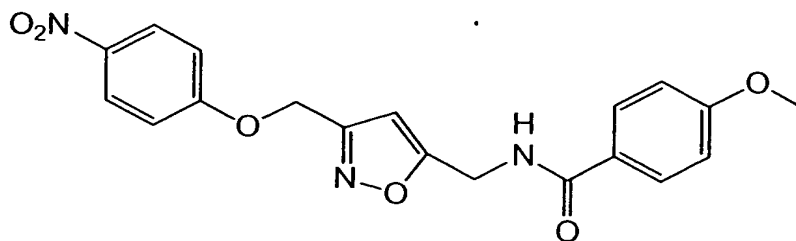


10

Melting point: 95 to 106°C.

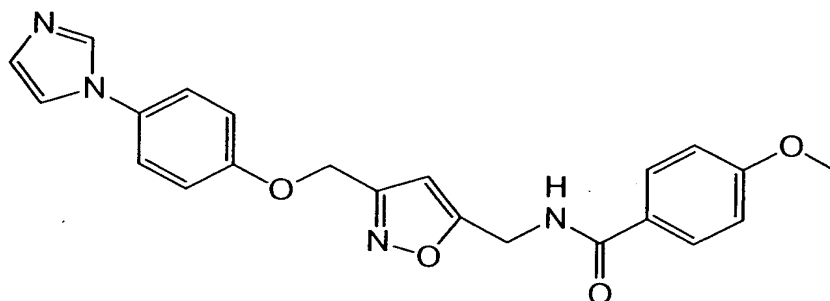
Example 17(114): Preparation of N-{{3-(4-nitrophenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-60):

15



Melting point: 151 to 154°C.

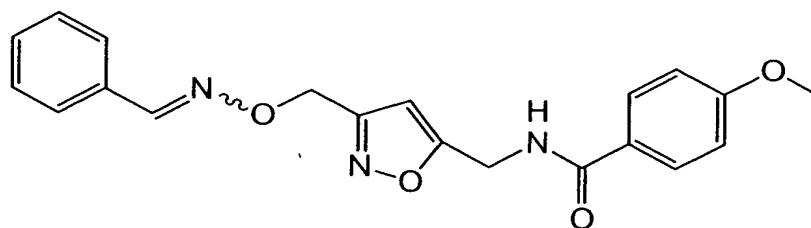
Example 17(115): Preparation of N-([3-(imidazol-2-ylphenyl)oxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide  
5 (Compound number IV-72):



Melting point: 138 to 142°C;

NMR :  $\delta$  3.80(3H, s), 4.70(2H, d), 5.08(2H, s), 6.32(1H, s),  
10 6.78-7.93(12H, m)

Example 17(116): Preparation of {3-[(4-methoxyphenyl)carbonylaminomethyl]-5-isoxazolyl}methoxyiminobenzyl (Compound number IV-74):



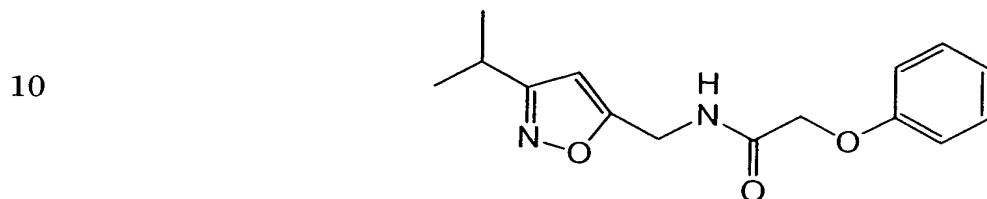
15 Property: amorphous;

NMR :  $\delta$  3.76(3H, s), 4.65(2H, d); 5.18(2H, s), 5.32(1H, s),  
6.25(1H, s), 6.73-8.09(11H, m)

#### Example 18

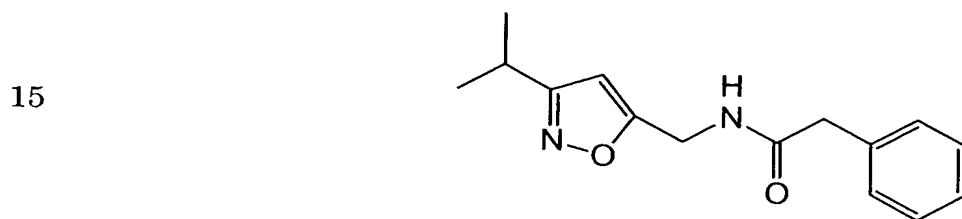
According to the methods described in Examples 1 to 16,  
5 the compounds in the following Examples 18(1) to 18(173)  
were obtained.

Example 18(1): Preparation of N-[(3-isopropyl-5-  
isoxazolyl)methyl]phenoxyacetamide (Compound number I-62):



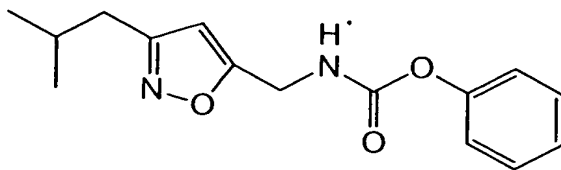
Melting point: 80 to 86°C.

Example 18(2): Preparation of N-[(3-isopropyl-5-  
isoxazolyl)methyl]phenylacetamide (Compound number I-63):



Melting point: 54 to 66°C.

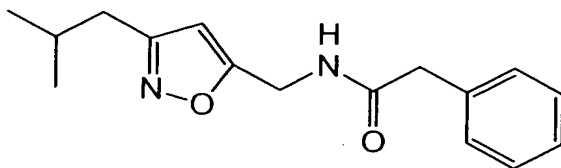
Example 18(3): Preparation of phenyl [(3-isobutyl-5-  
isoxazolyl)methyl] carbamate (Compound number I-8):



Property: oily;

NMR :  $\delta$  0.91(3H, s), 0.99(3H, s), 1.72-2.22(1H, m), 2.52(2H, d), 4.52(2H, d), 5.47(1H, br), 6.06(1H, s), 7.02-7.48(5H, m)

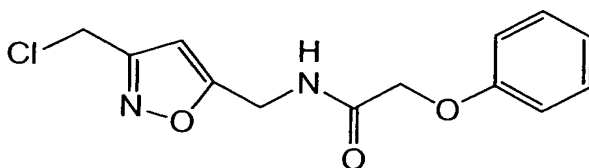
5 Example 18(4): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]phenylacetamide (Compound number I-64):



Property: oily;

10 NMR :  $\delta$  0.88(3H, s), 0.93(3H, s), 1.72-2.14(1H, m), 2.48(2H, d), 3.59(2H, s), 4.48(2H, d), 5.88(1H, s), 7.12-7.44(5H, m)

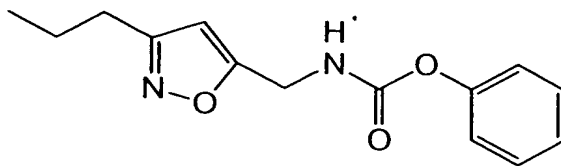
Example 18(5): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]phenoxyacetamide (Compound number I-65):



Melting point: 89 to 99°C.

15

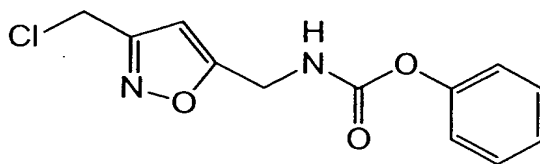
Example 18(6): Preparation of phenyl [(3-propyl-5-isoxazolyl)methyl] carbamate (Compound number I-3):



Property: oily;

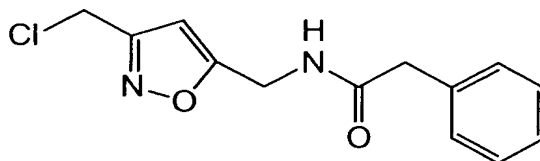
NMR :  $\delta$  0.92(3H, t), 1.43-1.88(2H, m), 2.60(2H, t), 4.48(2H, d), 5.89(1H, br), 6.07(1H, s), 7.00-7.50(5H, m)

- 5 Example 18(7): Preparation of phenyl [(3-chloromethyl-5-isoxazolyl)methyl] carbamate (Compound number I-66):



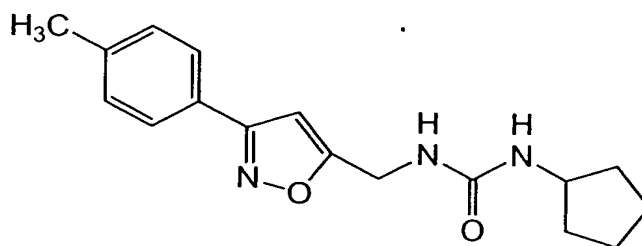
Melting point: 79 to 81°C.

- 10 Example 18(8): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]phenylacetamide (Compound number I-67):



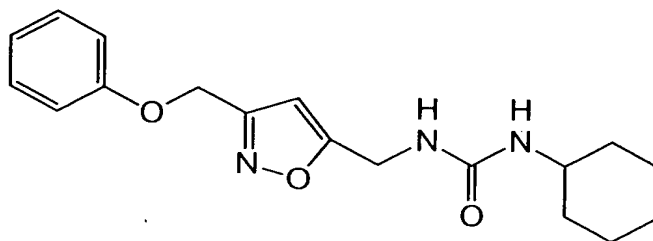
Melting point: 102 to 103°C.

- 15 Example 18(9): Preparation of 3-[3-(4-tolyl)-5-isoxazolyl]methyl-1-cyclopentylurea (Compound number II-10):



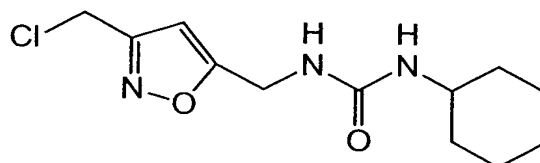
Melting point: 163 to 164°C.

Example 18(10): Preparation of 3-(3-phenoxyethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-104):



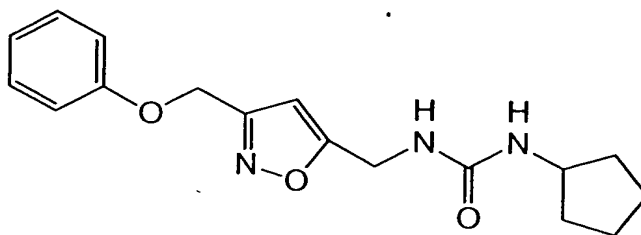
Melting point: 137 to 138°C.

Example 18(11): Preparation of 3-(3-chloromethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-105):



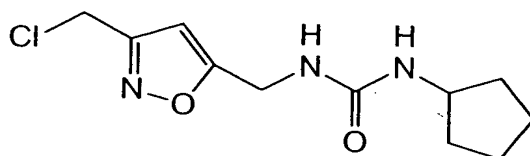
Melting point: 96 to 107°C.

Example 18(12): Preparation of 3-(3-phenoxyethyl-5-isoxazolyl)methyl-1-cyclopentylurea (Compound number II-106):



NMR :  $\delta$  1.10-2.20 (8H, m), 3.74-4.12 (1H, m), 4.42 (2H, d), 5.08 (2H, s), 5.13 (1H, br), 5.60 (1H, br), 6.28 (1H, s), 6.82-7.40 (5H, m)

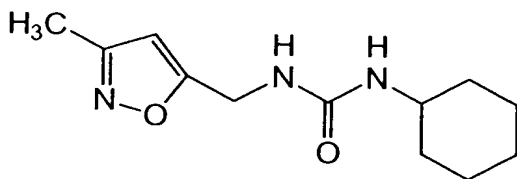
- 5 Example 18(13): Preparation of 3-(3-chloromethyl-5-isoxazolyl)methyl-1-cyclopentylurea (Compound number II-107):



Property: oily;

- 10 NMR :  $\delta$  0.72-2.20 (8H, m), 3.80-4.20 (1H, m), 4.50 (2H, d), 4.53 (2H, s), 4.40-5.35 (2H, m), 6.28 (1H, s)

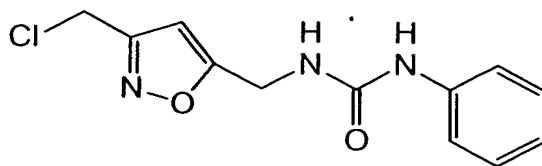
Example 18(14): Preparation of 3-(3-methyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-108):



- 15 Melting point: 119 to 122°C.

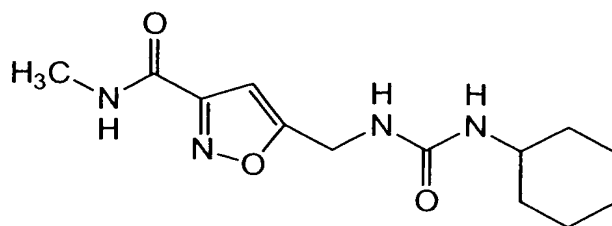
Example 18(15): Preparation of 3-(3-chloromethyl-5-isoxazolyl)methyl-1-phenylurea (Compound number II-109):





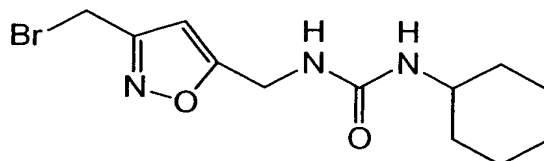
Melting point: 144 to 146°C.

Example 18(16): Preparation of 3-(3-methylaminocarbonyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-110):



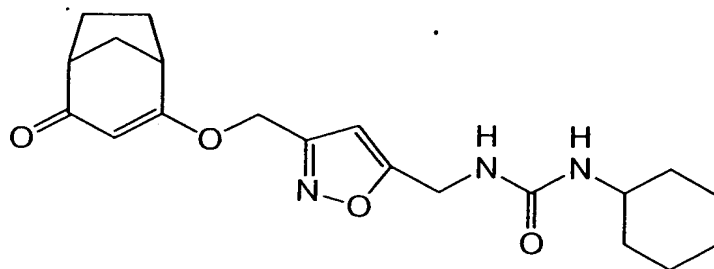
Melting point: 197 to 198°C.

Example 18(17): Preparation of 3-(3-bromomethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-111):



Melting point: 137 to 139°C.

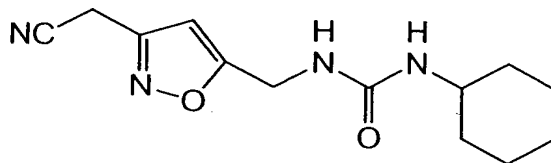
Example 18(18): Preparation of 3-[3-(bicyclo[3.2.1]oct-2-en-4-on-2-yl)oxymethyl-5-isoxazolyl]methyl-1-cyclohexylurea (Compound number II-112):



Property: gummy;

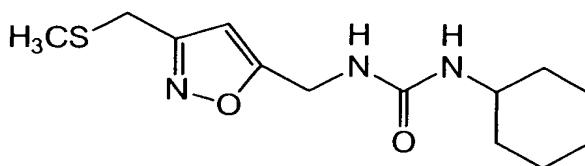
NMR :  $\delta$  0.60-2.40 (16H, m), 2.90 (2H, m), 3.50 (1H, m), 4.50 (2H, d), 4.85 (2H, s), 5.00 (1H, br), 5.10 (1H, s), 5.50 (1H, b), 6.25 (1H, s)

Example 18(19): Preparation of 3-(3-cyanomethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-113):



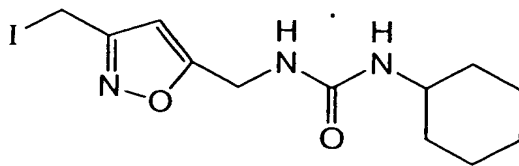
Melting point: 140 to 150°C.

Example 18(20): Preparation of 3-(3-methylthiomethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-114):



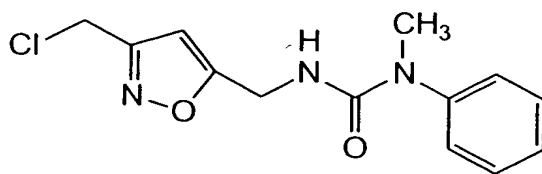
Melting point: 123 to 130°C.

Example 18(21): Preparation of 3-(3-iodomethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-115):



Melting point: 136 to 139°C.

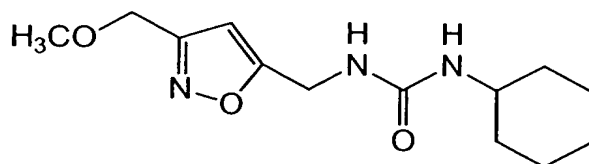
Example 18(22): Preparation of 3-[(3-chloromethyl-5-isoxazolyl)methyl]-1-methyl-1-phenylurea (Compound number II-116):



Property: oily;

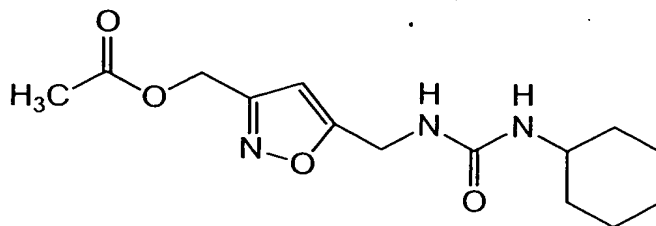
NMR :  $\delta$  3.25(3H,s), 4.50(2H,d), 4.55(2H,s), 4.70-4.90(1H,brs), 6.25(1H,s), 7.10-7.60(5H,m)

Example 18(23): Preparation of 3-(3-methoxymethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-117):



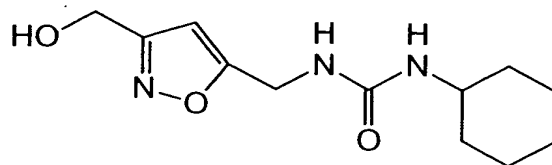
Melting point: 98 to 99°C.

Example 18(24): Preparation of 3-(3-acetyloxymethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-118):



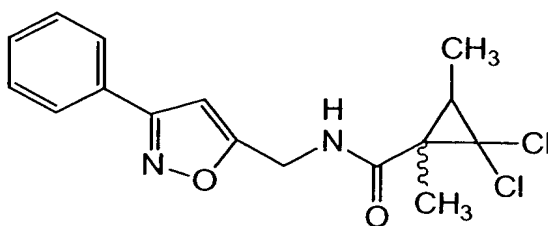
Melting point: 89 to 100°C.

Example 18(25): Preparation of 3-(3-hydroxymethyl-5-isoxazolyl)methyl-1-cyclohexylurea (Compound number II-119):



Melting point: 114 to 116°C.

Example 18(26): Preparation of N-[(3-phenyl-5-isoxazolyl)methyl]-2,2-dichloro-1,3-dimethylcyclopropanecarboxamide (Compound number III-212):

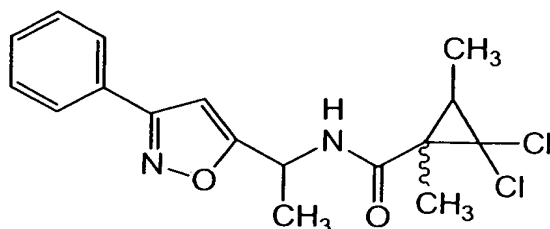


Property: oily;

NMR :  $\delta$  1.07-2.50 (7H, m), 4.65 (2H, d), 4.61 (1H, br), 6.52 (1H, s), 7.32-7.87 (5H, m)

Example 18(27): Preparation of N-[1-(3-phenyl-5-isoxazolyl)ethyl]-2,2-dichloro-1,3-

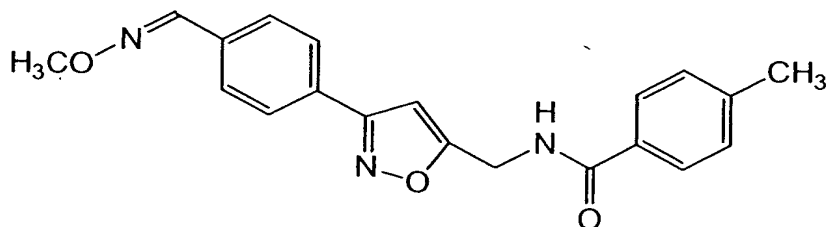
dimethylcyclopropanecarboxamide (Compound number III-213):



Property: gummy;

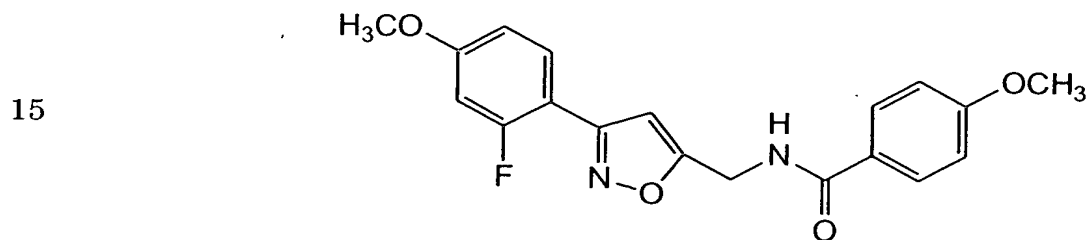
5 NMR :  $\delta$  1.07-2.47(10H, m), 5.21-5.65(1H, m), 6.02-6.35(1H, br), 6.41(0.5H, s), 6.48(0.5H, s), 7.32-7.89(5H, m)

Example 18(28): Preparation of N-([3-(methoxyiminomethyl)phenyl-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-214):



10 Melting point: 154 to 164°C.

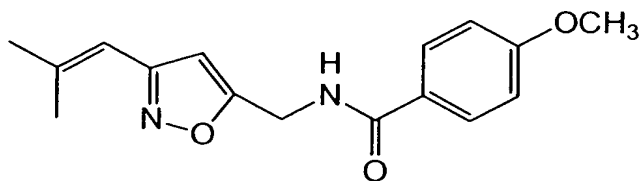
Example 18(29): Preparation of N-([3-(2-fluoro-4-methoxyphenyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-215):



15 Melting point: 129 to 131°C.

Example 18(30): Preparation of N-([3-(2-methyl-1-propenyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-216):

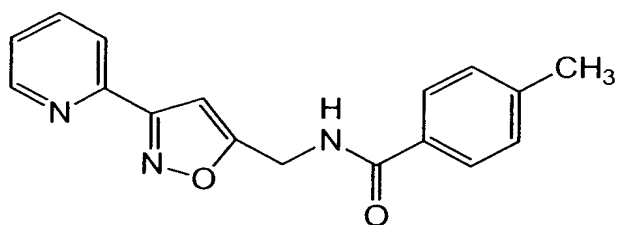
5



Melting point: 74 to 85°C.

Example 18(31): Preparation of N-([3-(2-pyridyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-217):

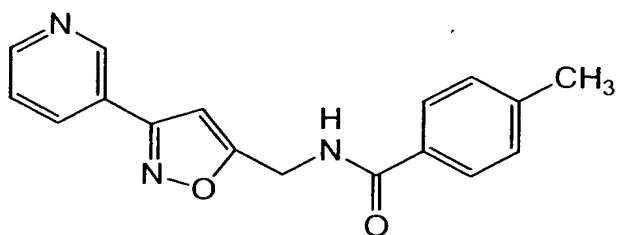
10



Melting point: 141 to 142°C.

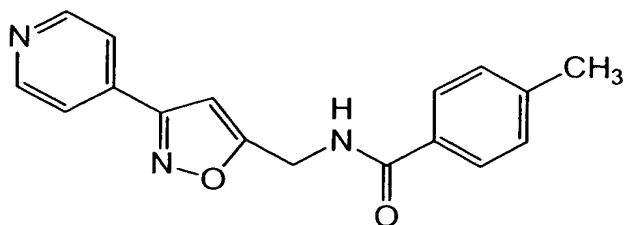
Example 18(32): Preparation of N-([3-(3-pyridyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-218):

15



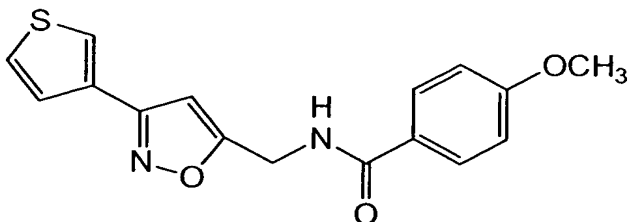
Melting point: 127 to 129°C.

Example 18(33): Preparation of N-([3-(4-pyridyl)-5-isoxazolyl]methyl)-4-methylbenzamide (Compound number III-  
5 219):



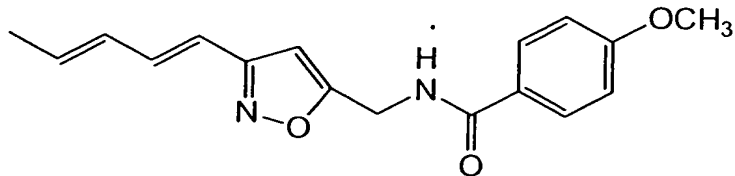
NMR :  $\delta$  2.40 (3H, s), 4.82 (2H, d), 6.62 (1H, s), 6.98 (1H, br),  
7.13-8.88 (8H, m)

Example 18(34): Preparation of N-([3-(3-thienyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-  
10 220):



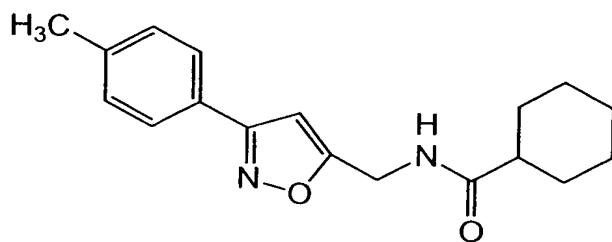
Melting point: 142 to 155°C.

15 Example 18(35): Preparation of N-([3-(1,3-pentadienyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-221):



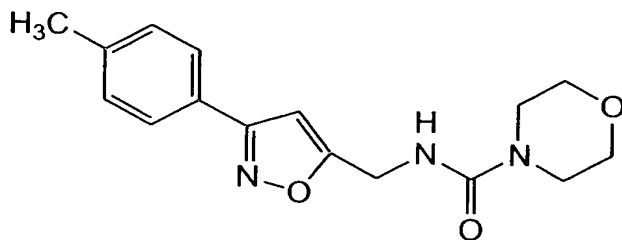
Melting point: 151 to 154°C.

Example 18(36): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)cyclohexanecarboxamide (Compound number III-222):



Melting point: 162 to 164°C.

10 Example 18(37): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-4-morpholinecarboxamide (Compound number III-204):



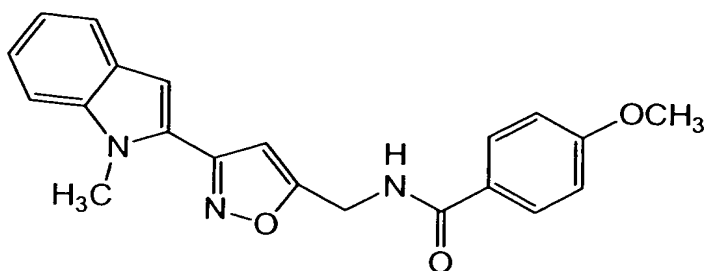
Melting point: 118 to 123°C.

15

Example 18(38): Preparation of N-([3-(1-methylindol-2-yl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-

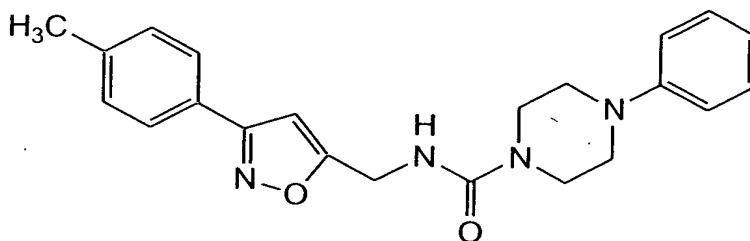


223):



Melting point: 110 to 114°C.

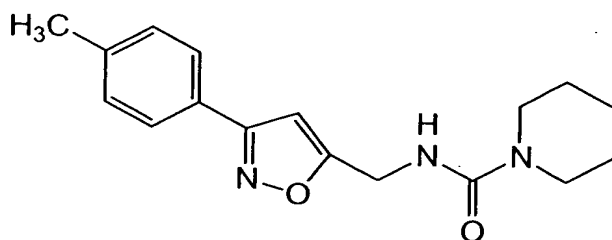
- 5 Example 18(39): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-4-phenylpiperazine-1-carboxamide (Compound number III-207):



Melting point: 177 to 179°C.

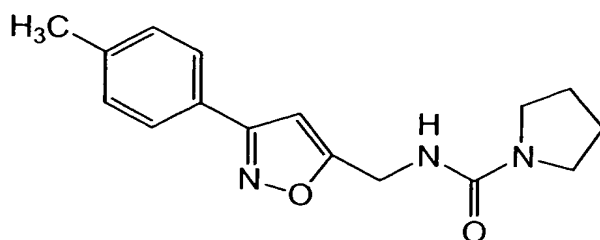
10

- Example 18(40): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-1-pyridinecarboxamide (Compound number III-203):



- 15 Melting point: 130 to 133°C.

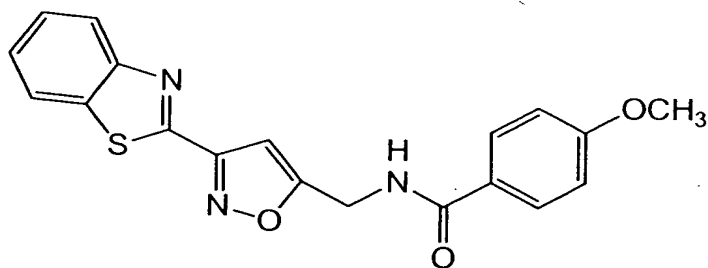
Example 18(41): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-1-pyrrolidinecarboxamide (Compound number III-205):



5 Melting point: 131 to 142°C.

Example 18(42): Preparation of N-([3-(benzo[d]thiazol-2-yl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-224):

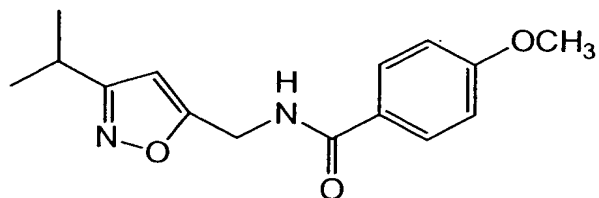
10



NMR :  $\delta$  3.85(3H, s), 4.74(2H, d), 6.72(1H, s), 6.69-8.34(4H, m), 9.42(1H, br)

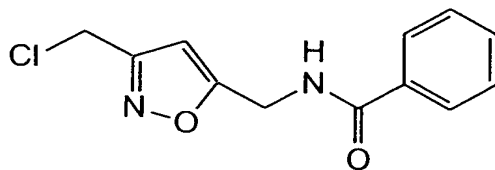
Example 18(43): Preparation of N-([3-isopropyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number III-225):

15



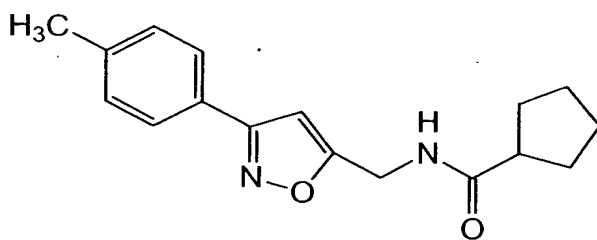
Melting point: 114 to 119°C.

Example 18(44): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]benzamide (Compound number III-226):



5 Melting point: 109 to 117°C.

Example 18(45): Preparation of N-[[3-(4-tolyl)-5-isoxazolyl]methyl]cyclopentanecarboxamide (Compound number III-227):

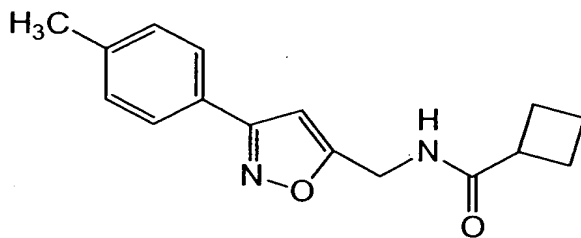


10

Melting point: 162 to 164°C.

Example 18(46): Preparation of N-[[3-(4-tolyl)-5-isoxazolyl]methyl]cyclobutanecarboxamide (Compound number III-228):

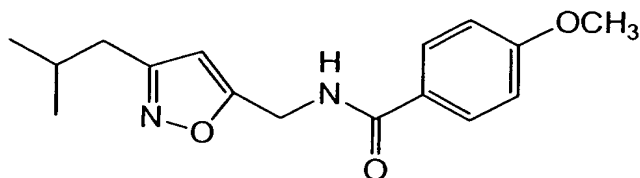
15



Melting point: 168 to 170°C.

Example 18(47): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-229):

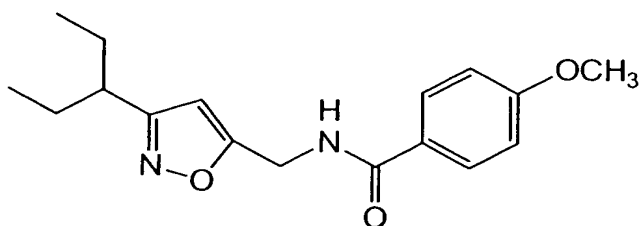
5



Melting point: 94 to 113°C.

Example 18(48): Preparation of N-[[3-(1-ethylpropyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number III-230):

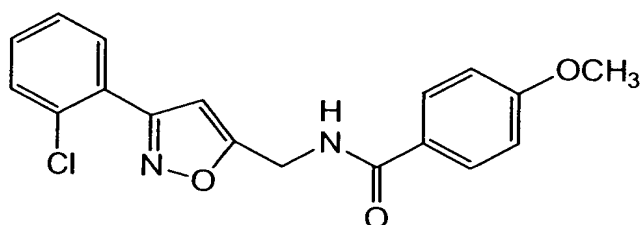
10



Melting point: 95 to 114°C.

Example 18(49): Preparation of N-[[3-(2-chlorophenyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number III-231):

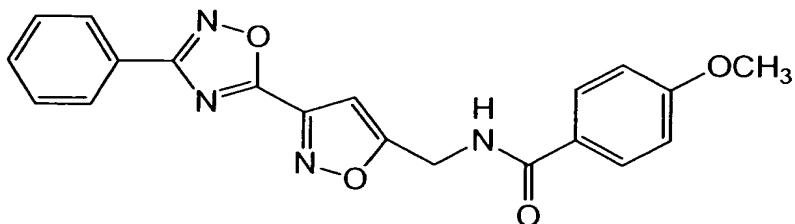
15



Melting point: 87 to 101°C.

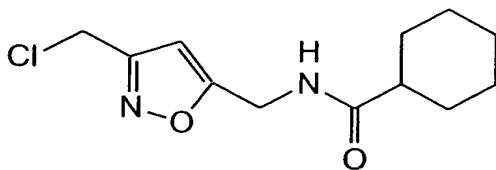
Example 18(50): Preparation of N-([3-(3-phenyl-1,2,4-oxadiazol-5-yl)-5-isoxazolyl]methyl)-4-methoxybenzamide

5 (Compound number III-232):



Melting point: 150 to 162°C.

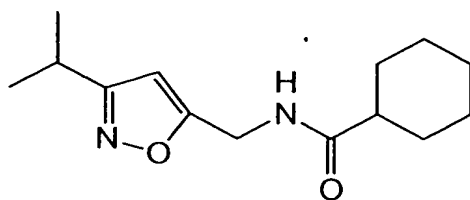
Example 18(51): Preparation of N-([3-chloromethyl-5-isoxazolyl]methyl)cyclohexanecarboxamide (Compound number III-233):



Property: paste;

15 NMR :  $\delta$  1.03-2.33(11H, m), 4.52(2H, s), 4.52(2H, d), 6.03(1H, br), 6.26(1H, s)

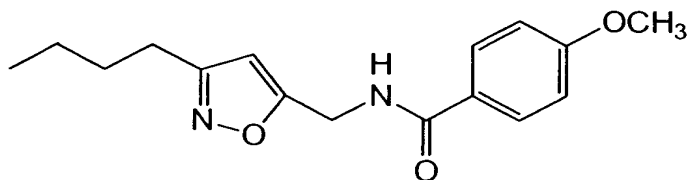
Example 18(52): Preparation of N-([3-isopropyl-5-isoxazolyl]methyl)cyclohexanecarboxamide (Compound number III-234):



Property: oily;

NMR :  $\delta$  0.87-2.27(10H, m), 1.20(3H, s), 1.30(3H, s), 2.82-3.23(1H, m), 4.50(2H, d), 5.92(1H, br), 6.00(1H, s)

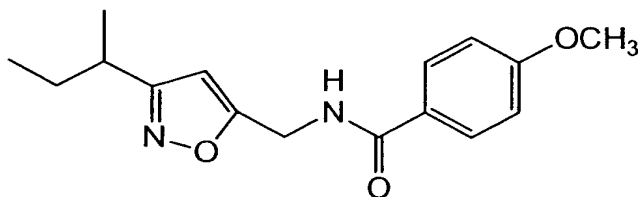
5 Example 18(53): Preparation of N-[(3-butyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-235):



Melting point: 101 to 104°C.

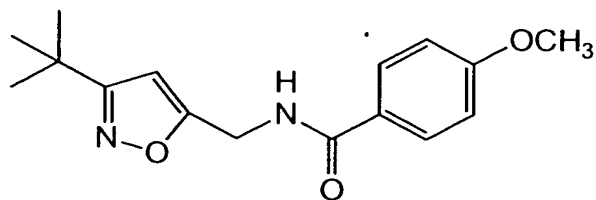
10

Example 18(54): Preparation of N-[(3-sec-butyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-236):



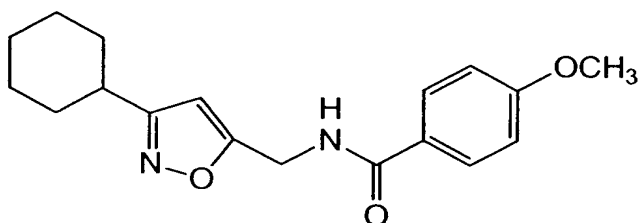
15 Melting point: 70 to 80°C.

Example 18(55): Preparation of N-[(3-tert-butyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-237):



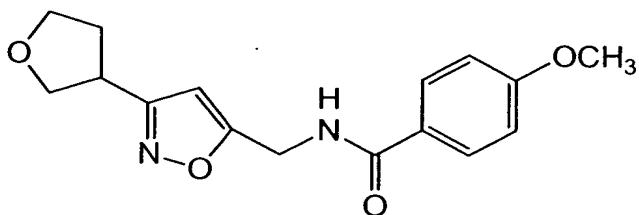
Melting point: 125 to 132°C.

Example 18(56): Preparation of N-[(3-cyclohexyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-238):



Melting point: 93 to 107°C.

10 Example 18(57): Preparation of N-[(3-(oxolan-3-yl)-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-239):

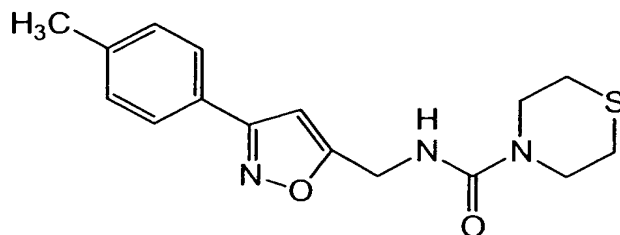


Melting point: 113 to 117°C.

15

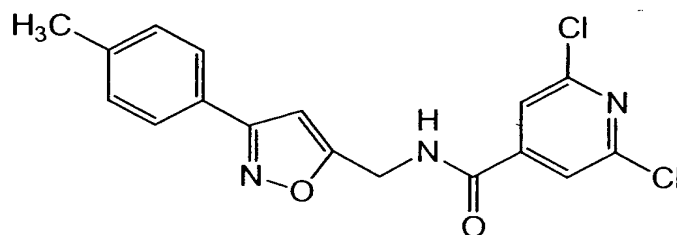
Example 18(58): Preparation of N-[(3-(4-tolyl)-5-isoxazolyl)methyl]-1,4-thiazinane-4-carboxamide (Compound

number III-206):



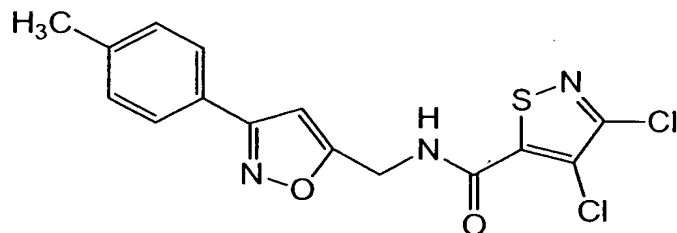
Melting point: 141 to 145°C.

- 5 Example 18(59): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-2,6-dichloropyridine-4-carboxamide  
(Compound number III-240):



Melting point: 149 to 151°C.

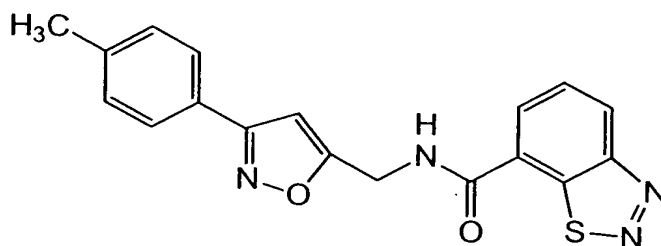
- 10 Example 18(60): Preparation of N-([3-(4-tolyl)-5-isoxazolyl]methyl)-3,4-dichloroisothiazole-5-carboxamide  
(Compound number III-241):



- 15 Melting point: 155 to 157°C.



Example 18(61): Preparation of N-([3-(4-tolyl)-5-isoxazolyl)methyl]-benzo[d][1,2,3]thiazole-7-carboxamide  
(Compound number III-242):

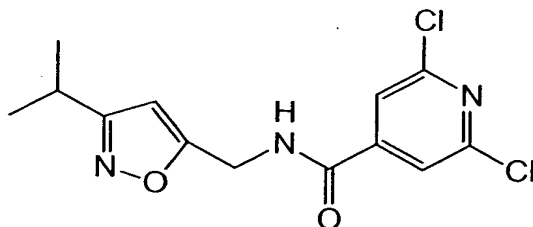


5

Melting point: 178 to 192°C.

Example 18(62): Preparation of N-[(3-isopropyl-5-isoxazolyl)methyl]-2,6-dichloropyridine-4-carboxamide

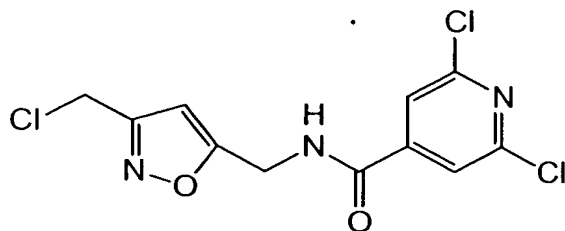
10 (Compound number III-243):



Property: oily;

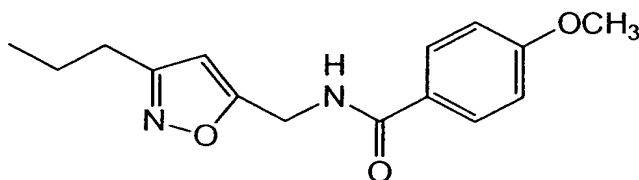
NMR :  $\delta$  1.19(3H, s), 1.28(3H, s), 2.73-3.22(1H, m), 4.70(2H, d), 6.13(1H, s), 7.68(2H, s), 8.00(1H, br)

15 Example 18(63): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,6-dichloropyridine-4-carboxamide  
(Compound number III-244):



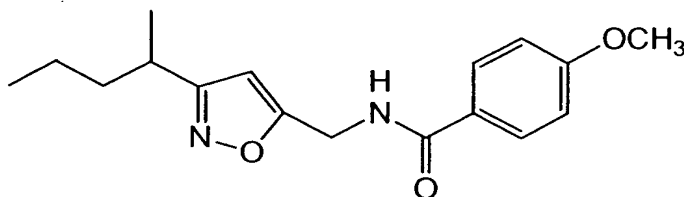
NMR :  $\delta$  4.54(2H, s), 4.73(2H, d), 6.34(1H, s), 6.81(1H, br), 7.59(2H, s)

Example 18(64): Preparation of N-[(3-propyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-245):



Melting point: 97 to 102°C.

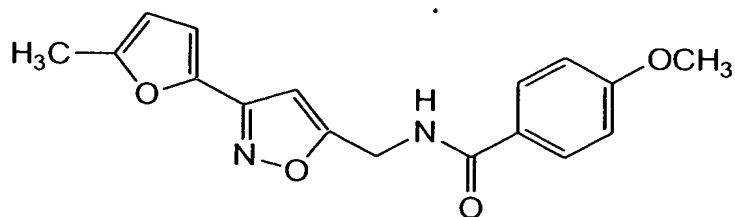
10 Example 18(65): Preparation of N-[(3-sec-pentyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-246):



Melting point: 72 to 76°C.

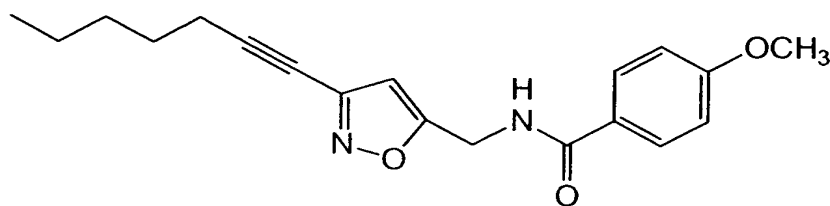
15

Example 18(66): Preparation of N-[[3-(2-methylfuran-5-yl)-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-247):



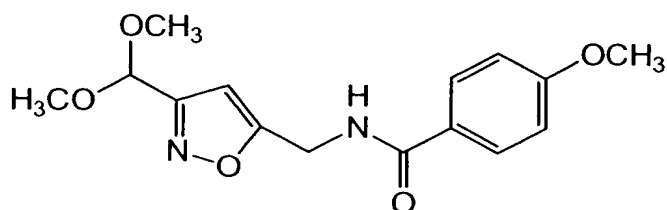
Melting point: 107 to 109°C.

Example 18(67): Preparation of N-[[3-(1-heptynyl)-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number III-248):



Melting point: 96 to 102°C.

10 Example 18(68): Preparation of N-[(3-dimethoxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-249):

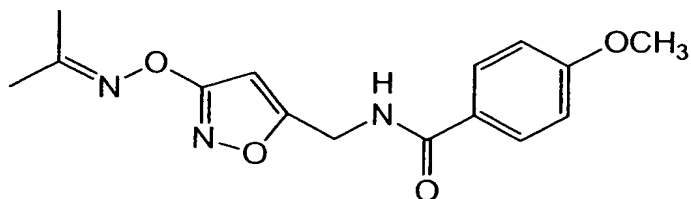


Property: paste;

15 NMR :  $\delta$  3.36(6H, s), 3.83(3H, s), 4.73(2H, d), 5.40(1H, s), 6.28(1H, s), 6.73(1H, br), 6.88(2H, d), 7.74(2H, d)

Example 18(69): Preparation of N-[(3-isopropyliminoxy-5-

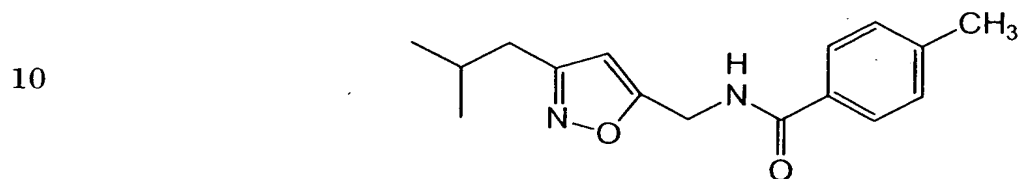
isoxazolyl)methyl]-4-methoxybenzamide (Compound number III-250):



Property: paste;

5 NMR :  $\delta$  1.81(6H, s), 3.83(3H, s), 4.68(2H, d), 5.02(2H, s), 6.22(1H, s), 6.86(2H, d), 7.74(2H, d)

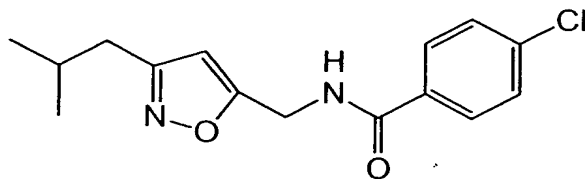
Example 18(70): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number III-251):



Melting point: 131 to 134°C.

Example 18(71): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-252):

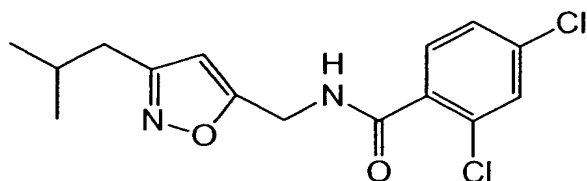
15



Melting point: 122 to 124°C.

Example 18(72): Preparation of N-[(3-isobutyl-5-

isoxazolyl)methyl]-2,4-dichlorobenzamide (Compound number  
III-253):

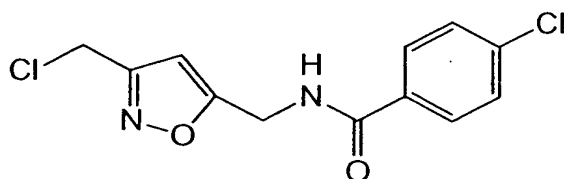


Property: oily;

5 NMR :  $\delta$  0.88(3H, s), 0.96(3H, s), 1.76-2.13(1H, m), 2.51(2H, d), 4.71(2H, d), 6.09(1H, s), 6.84(1H, br), 7.16-7.70(3H, m)

Example 18(73): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-254):

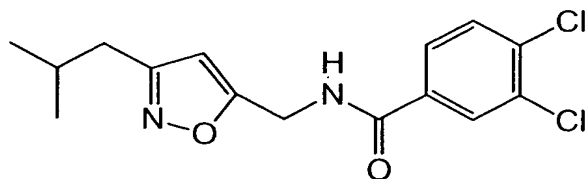
10



Melting point: 145 to 147°C.

Example 18(74): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-3,4-dichlorobenzamide (Compound number

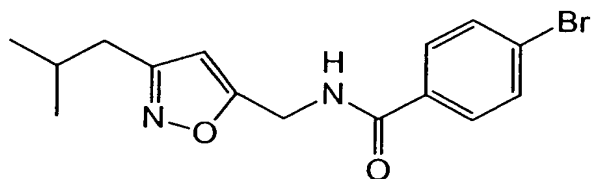
15 III-255):



Melting point: 68 to 74°C.

Example 18(75): Preparation of N-[(3-isobutyl-5-

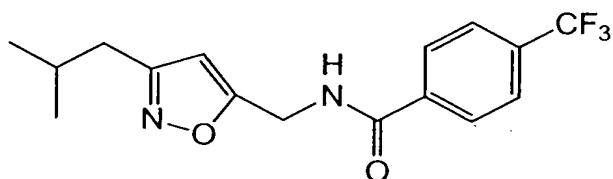
isoxazolyl)methyl]-4-bromobenzamide (Compound number III-256):



Melting point: 124 to 125°C.

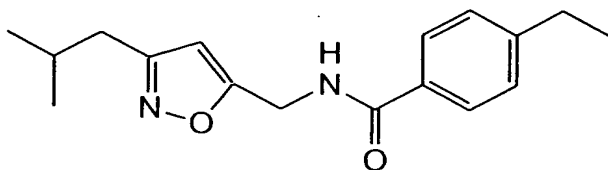
5

Example 18(76): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-4-trifluoromethylbenzamide (Compound number III-257):



10 Melting point: 111 to 112°C.

Example 18(77): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-4-ethylbenzamide (Compound number III-258):

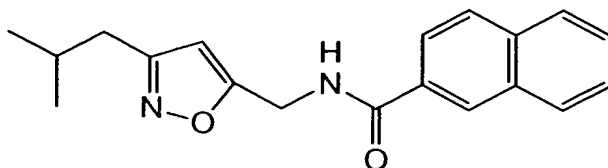


15

Melting point: 127 to 128°C.

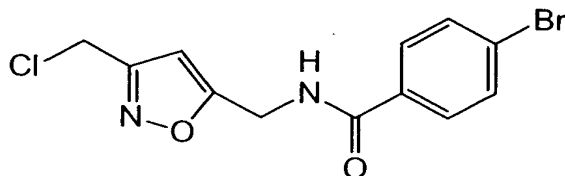
Example 18(78): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-2-naphthalenecarboxamide (Compound number

III-259):



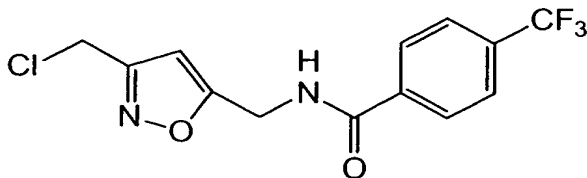
Melting point: 109 to 118°C.

- 5 Example 18(79): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-bromobenzamide (Compound number III-260):



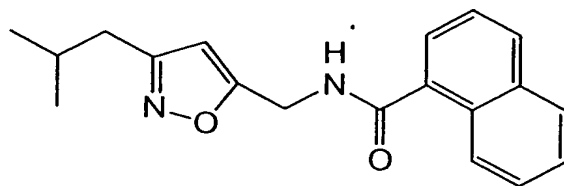
10 NMR :  $\delta$  4.55(2H, s), 4.70(2H, d), 6.35(1H, s), 7.64(4H, d), 7.97(1H, br)

- Example 18(80): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-trifluoromethylbenzamide (Compound number III-261):



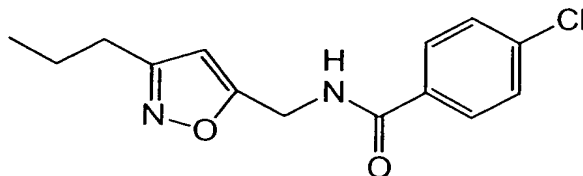
15 Melting point: 145 to 147°C.

- Example 18(81): Preparation of N-[(3-isobutyl-5-isoxazolyl)methyl]-1-naphthalenecarboxamide (Compound number III-262):



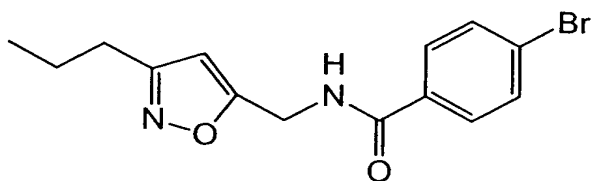
Melting point: 89 to 92°C.

Example 18(82): Preparation of N-[(3-propyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-263):



Melting point: 133 to 134°C.

10 Example 18(83): Preparation of N-[(3-propyl-5-isoxazolyl)methyl]-4-bromobenzamide (Compound number III-264):

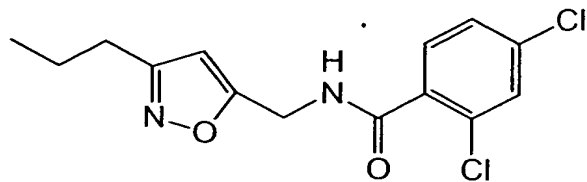


Melting point: 132 to 133°C.

15

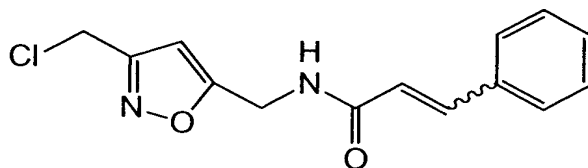
Example 18(84): Preparation of N-[(3-propyl-5-isoxazolyl)methyl]-2,4-dichlorobenzamide (Compound number III-265):





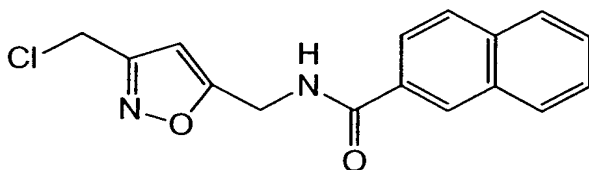
Melting point: 67 to 71°C.

Example 18(85): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-β-phenylacrylamide (Compound number III-266):



NMR :  $\delta$  4.52(2H, s), 4.67(2H, d), 6.30(1H, s), 6.35(0.5H, s), 6.52(0.5H, s), 6.80(1H, br), 7.13-7.75(6H, m)

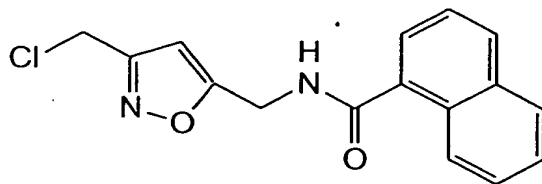
10 Example 18(86): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-naphthalenecarboxamide (Compound number III-267):



Melting point: 106 to 119°C.

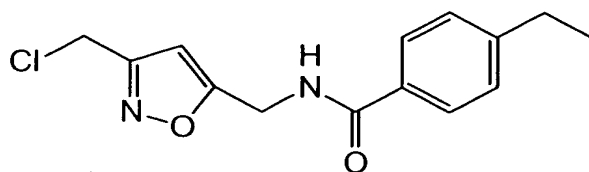
15

Example 18(87): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-1-naphthalenecarboxamide (Compound number III-268):



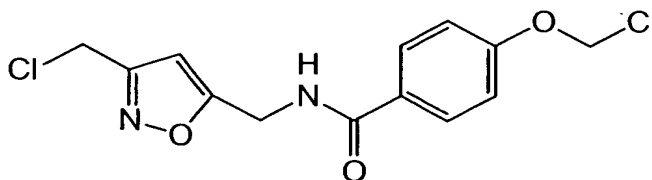
Melting point: 120 to 123°C.

Example 18(88): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-ethylbenzamide (Compound number III-269):



NMR :  $\delta$  1.22(3H, t), 2.65(2H, q), 4.51(2H, s), 4.72(2H, d), 6.31(1H, s), 6.72(1H, br), 7.20(2H, d), 7.71(2H, d)

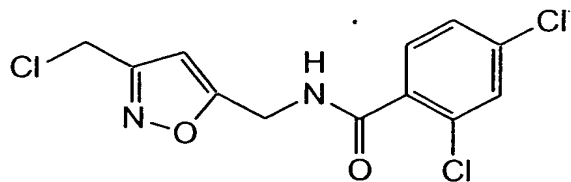
10 Example 18(89): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-chloromethoxybenzamide (Compound number III-270):



Melting point: 85 to 88°C.

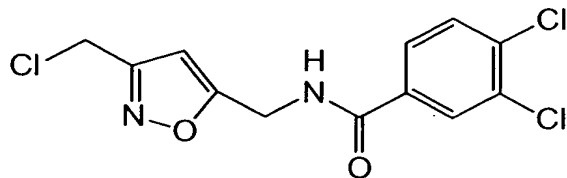
15

Example 18(90): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,4-dichlorobenzamide (Compound number III-271):



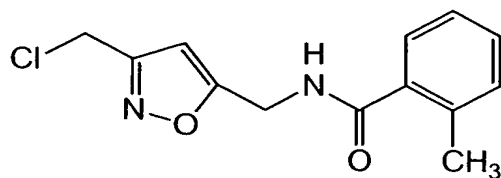
Melting point: 90 to 91°C.

Example 18(91): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-3,4-dichlorobenzamide (Compound number III-272):



Melting point: 119 to 121°C.

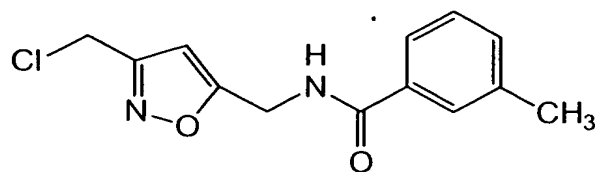
10 Example 18(92): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-methylbenzamide (Compound number III-273):



Melting point: 76 to 83°C.

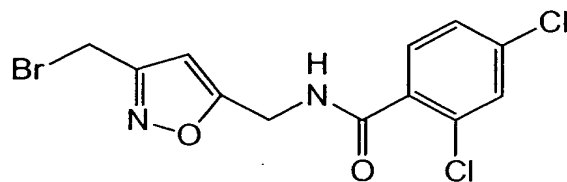
15

Example 18(93): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-3-methylbenzamide (Compound number III-274):



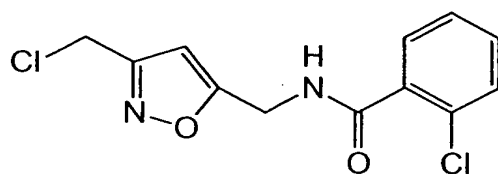
Melting point: 87 to 93°C.

Example 18(94): Preparation of N-[(3-bromomethyl-5-isoxazolyl)methyl]-2,4-dichlorobenzamide (Compound number III-275):



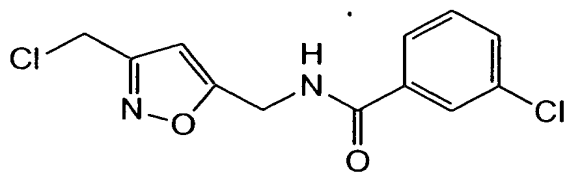
Melting point: 108 to 109°C.

10 Example 18(95): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-chlorobenzamide (Compound number III-276):



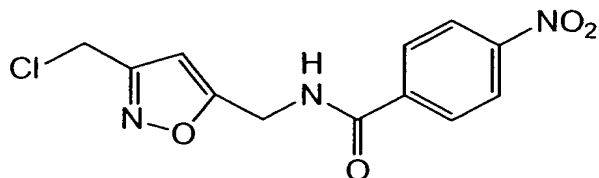
Melting point: 108 to 109°C.

15 Example 18(96): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-3-chlorobenzamide (Compound number III-277):



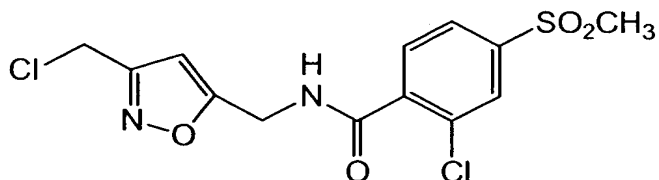
Melting point: 102 to 104°C.

Example 18(97): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-nitrobenzamide (Compound number III-278):



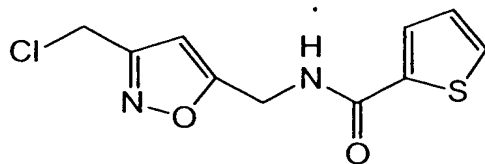
Melting point: 125 to 128°C.

10 Example 18(98): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-chloro-4-methanesulfonylbenzamide (Compound number III-279):



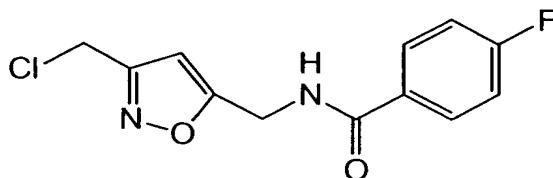
Melting point: 107 to 110°C.

15 Example 18(99): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-thiophenecarboxamide (Compound number III-280):



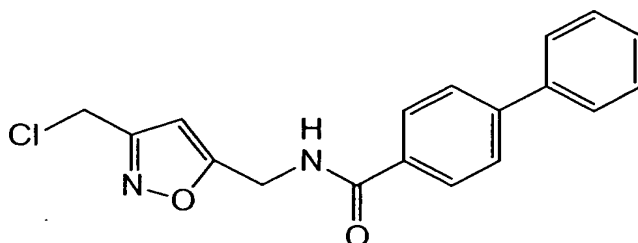
Melting point: 117 to 125°C.

Example 18(100): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-fluorobenzamide (Compound number III-281):



Melting point: 116 to 120°C.

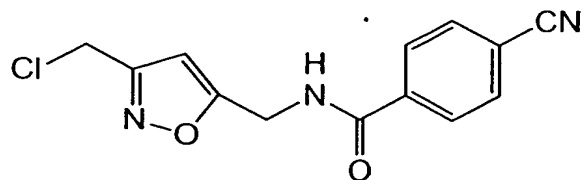
10 Example 18(101): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-phenylbenzamide (Compound number III-282):



Melting point: 129 to 139°C.

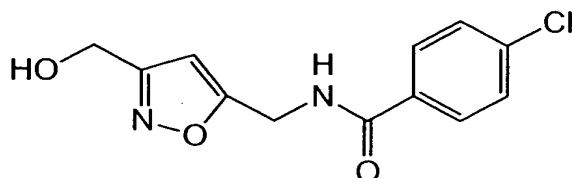
15

Example 18(102): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-cyanobenzamide (Compound number III-283):



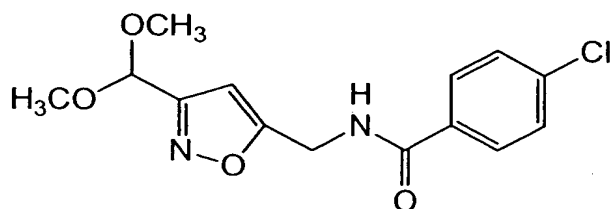
Melting point: 124 to 126°C.

Example 18(103): Preparation of N-[(3-hydroxymethyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-284):



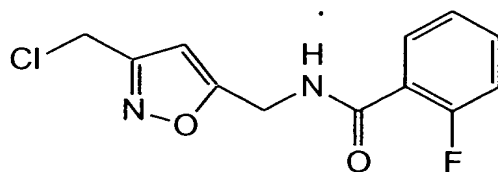
Melting point: 135 to 139°C.

10 Example 18(104): Preparation of N-[(3-dimethoxymethyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-285):



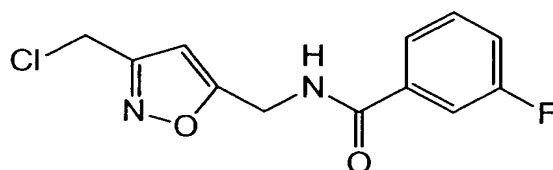
Melting point: 84 to 86°C.

15 Example 18(105): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-fluorobenzamide (Compound number III-286):



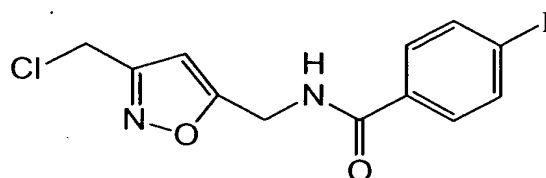
Melting point: 95 to 97°C.

Example 18(106): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-3-fluorobenzamide (Compound number III-287):



Melting point: 117 to 118°C.

10 Example 18(107): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-iodobenzamide (Compound number III-288):

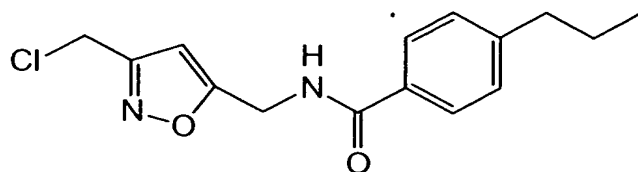


Melting point: 169 to 172°C.

15

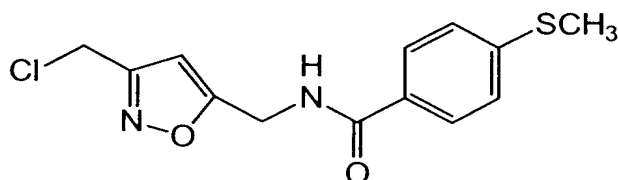
Example 18(108): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-propylbenzamide (Compound number III-289):





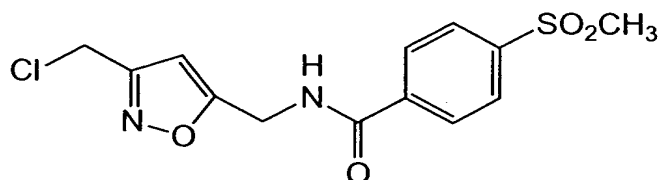
Melting point: 110 to 113°C.

Example 18(109): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-methylthiobenzamide (Compound number III-290):



Melting point: 147 to 149°C.

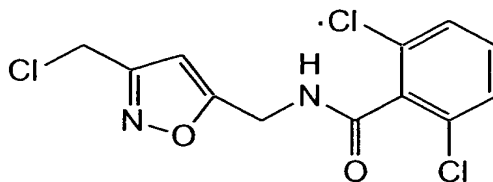
10 Example 18(110): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-methanesulfonylbenzamide (Compound number III-291):



Melting point: 134 to 135°C.

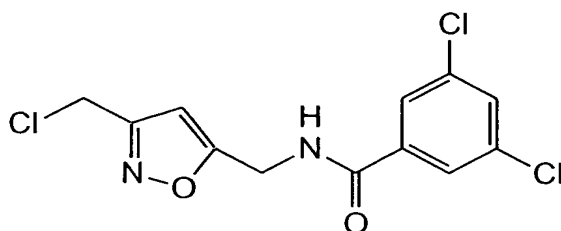
15

Example 18(111): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,6-dichlorobenzamide (Compound number III-292):



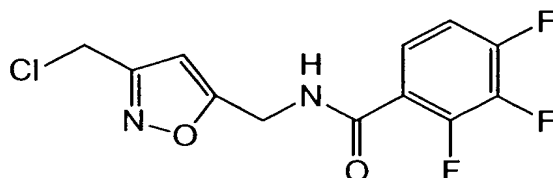
Melting point: 94 to 95°C.

Example 18(112): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-3,5-dichlorobenzamide (Compound number III-293):



Melting point: 136 to 137°C.

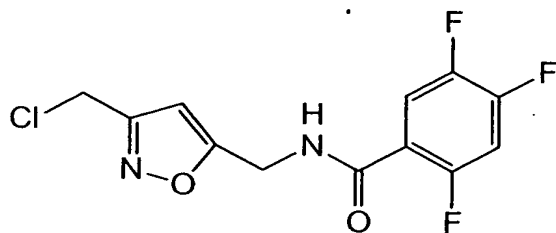
10 Example 18(113): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,3,4-trifluorobenzamide (Compound number III-294):



Melting point: 105 to 106°C.

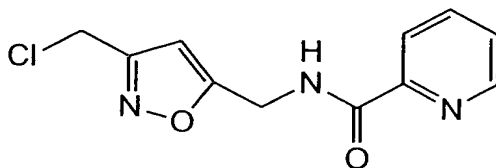
15

Example 18(114): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,4,5-trifluorobenzamide (Compound number III-295):



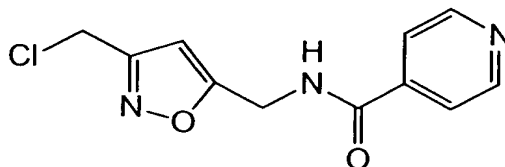
Melting point: 116 to 118°C.

Example 18(115): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2-pyridinecarboxamide (Compound number III-296):



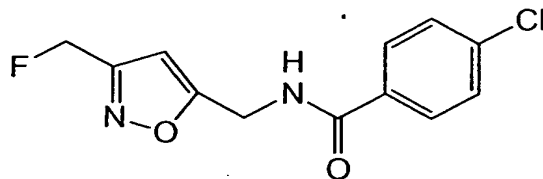
Melting point: 132 to 134°C.

10 Example 18(116): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-4-pyridinecarboxamide (Compound number III-297):



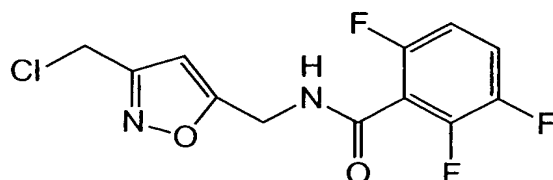
Melting point: 104 to 105°C.

15 Example 18(117): Preparation of N-[(3-fluoromethyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number III-298):



Melting point: 147 to 149°C.

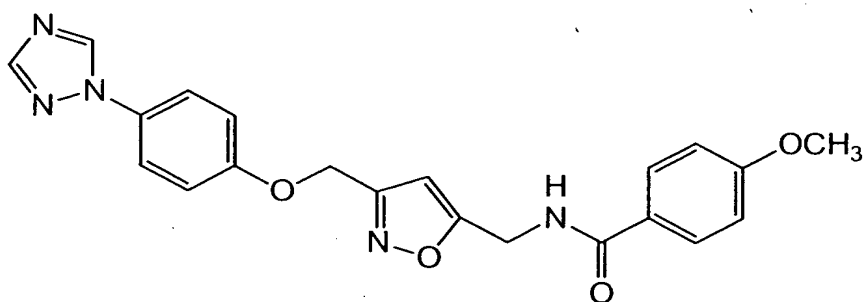
Example 18(118): Preparation of N-[(3-chloromethyl-5-isoxazolyl)methyl]-2,3,6-trifluorobenzamide (Compound number III-299):



Property: oily;

NMR :  $\delta$  4.50 (2H, s), 4.60 (2H, d), 6.25 (1H, s), 6.60-7.00 (1H, m),  
 10 7.00-7.20 (1H, m), 7.45-7.70 (1H, brs)

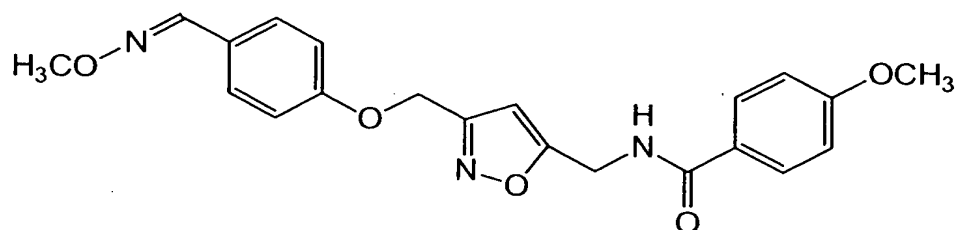
Example 18(119): Preparation of N-[[3-(1,2,4-triazol-1-yl)phenoxy)methyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-73):



15 NMR :  $\delta$  3.87 (3H, s), 4.72 (2H, d), 5.17 (2H, s), 6.35 (1H, s),  
 6.80-8.50 (10H, m), 9.08 (1H, s)

Example 18(120): Preparation of N-[[3-(4-

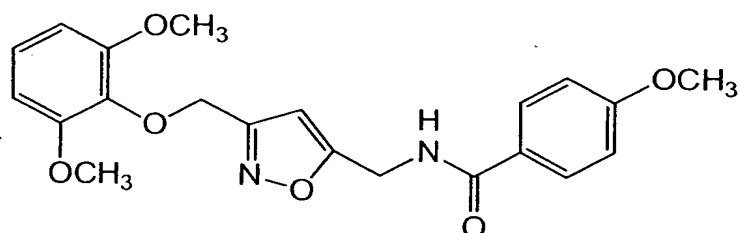
methoxyiminomethylphenyl)oxymethyl-5-isoxazolyl)methyl}-4-methoxybenzamide (Compound number IV-75):



Melting point: 143 to 151°C.

5

Example 18(121): Preparation of N-([3-(2,6-dimethoxyphenyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-76):

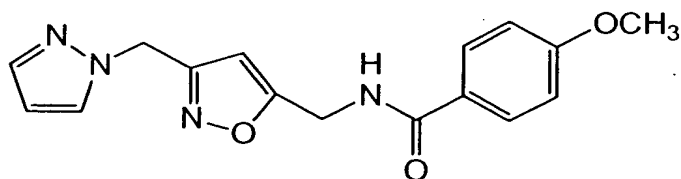


10 Property: paste;

NMR :  $\delta$  3.73(3H, s), 3.80(3H, s), 3.83(3H, s), 4.67(2H, d), 5.04(2H, s), 6.45(1H, s), 6.49-7.85(8H, m)

Example 18(122): Preparation of N-([3-(1,2-diazol-1-yl)methyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound

15 number IV-77):

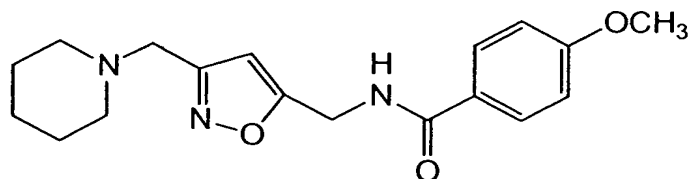


Property: paste;

NMR :  $\delta$  3.81(3H, s), 4.68(2H, s); 4.68(2H, d), 6.24(1H, s),  
6.28(1H, br), 6.70(1H, br), 6.77-7.90(6H, m)

Example 18(123): Preparation of N-[[3-(1-piperidyl)methyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-

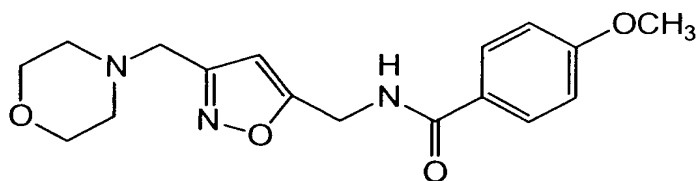
5 24):



Property: oily;

NMR :  $\delta$  3.51(2H, s), 3.84(3H, s), 4.73(2H, d), 6.27(1H, s),  
6.73(1H, br), 6.91(2H, d), 7.77(2H, d)

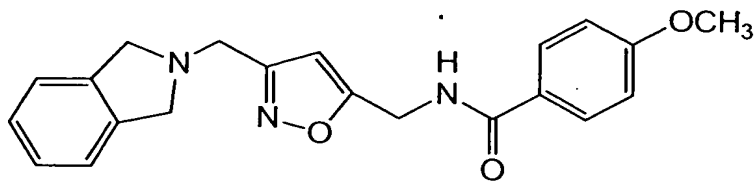
10 Example 18(124): Preparation of N-[(3-morpholinomethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-  
25):



Property: paste;

15 NMR :  $\delta$  2.32-2.52(4H, m), 3.52(2H, s), 3.56-3.73(4H, m),  
3.83(3H, s), 4.68(2H, d), 6.21(1H, s), 6.87(2H, d), 7.30(1H,  
br), 7.80(1H, d)

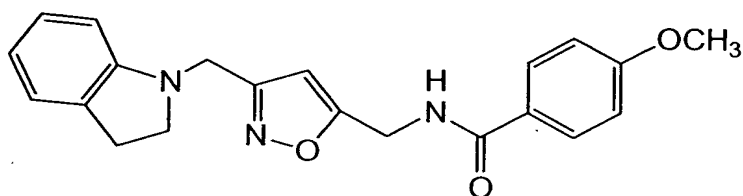
Example 18(125): Preparation of N-[[3-(1,3-dihydroindol-2-yl)methyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound  
20 number IV-78):



Property: paste;

NMR :  $\delta$  3.84(3H, s), 3.92(2H, s), 3.98(4H, s), 4.72(2H, d),  
 5 6.28(1H, s), 6.56(1H, br), 6.90(2H, d), 7.14(4H, s), 7.73(2H,  
 d)

Example 18(126): Preparation of N-([3-(2,3-dihydroindol-1-yl)methyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-79):

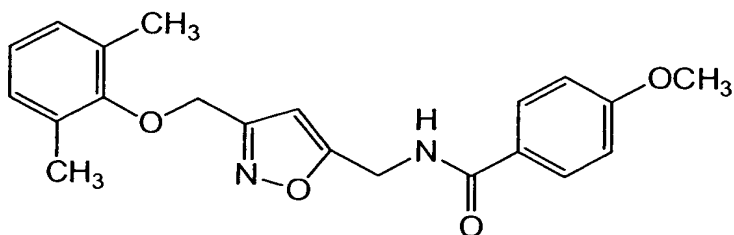


10 Property: paste;

NMR :  $\delta$  2.80-3.28(2H, m), 3.80(3H, s), 3.72-4.28(2H, m),  
 4.68(2H, d), 5.28(2H, s), 6.30(1H, s), 6.72-7.87(9H, m)

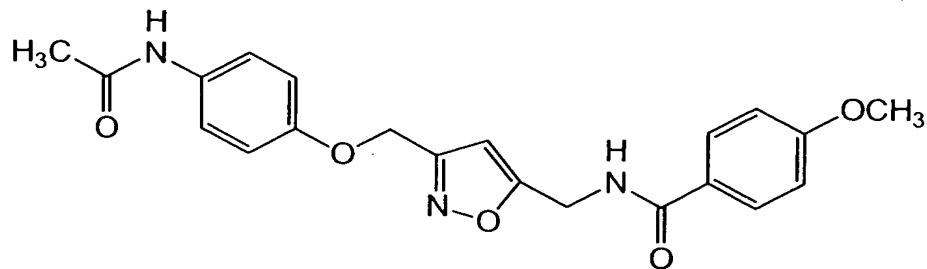
Example 18(127): Preparation of N-([3-(2,6-dimethylphenyl)oxymethyl-5-isoxazolyl]methyl)-4-

15 methoxybenzamide (Compound number IV-80):



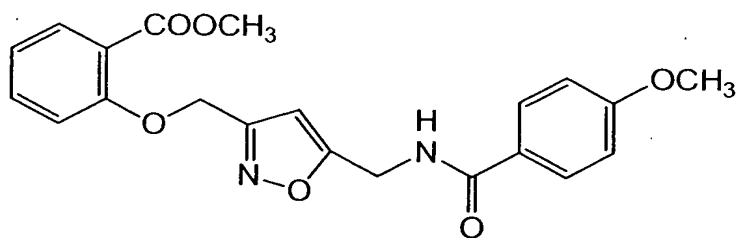
Melting point: 139 to 146°C.

Example 18(128): Preparation of N-{{3-(4-acetylamino-phenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-81):



5 Melting point: 183 to 188°C.

Example 18(129): Preparation of N-{{3-(2-methoxycarbonylphenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-82):



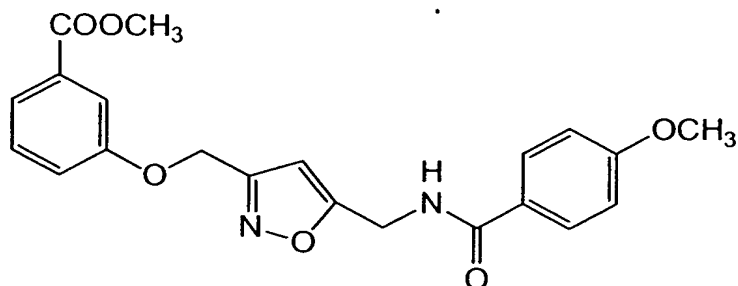
10

Melting point: 85 to 88°C.

Example 18(130): Preparation of N-{{3-(3-methoxycarbonylphenyl)oxymethyl-5-isoxazolyl}methyl}-4-methoxybenzamide (Compound number IV-83):

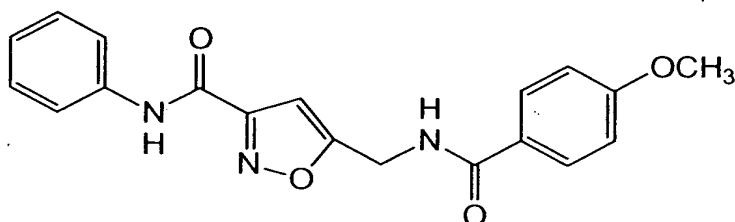
15





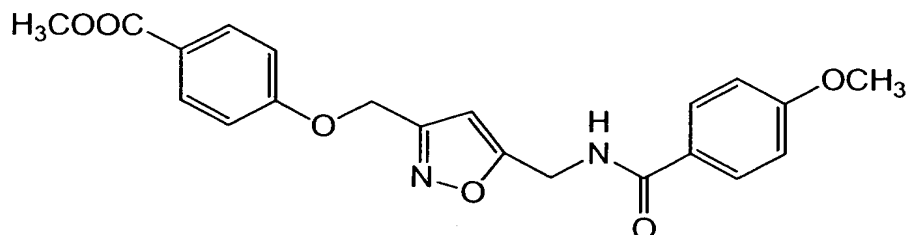
Melting point: 106 to 112°C.

Example 18(131): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-84):



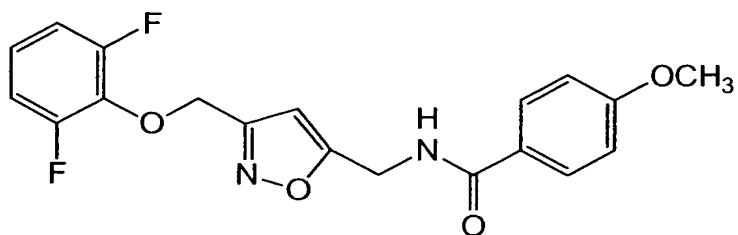
Melting point: 186°C.

10 Example 18(132): Preparation of N-[[3-(4-methoxycarbonylphenyl)oxymethyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-85):



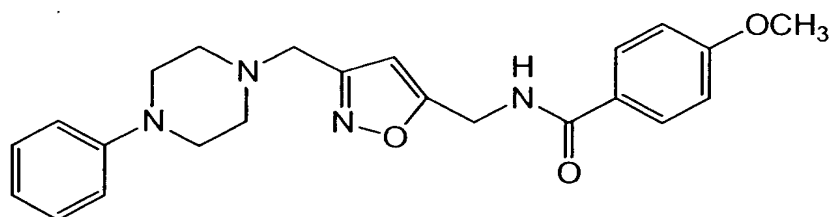
Melting point: 155 to 162°C.

Example 18(133): Preparation of N-[[3-(2,6-difluorophenyl)oxymethyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-86):



5 Melting point: 111 to 116°C.

Example 18(134): Preparation of N-[[3-(4-phenylpiperazinyl)methyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-28):

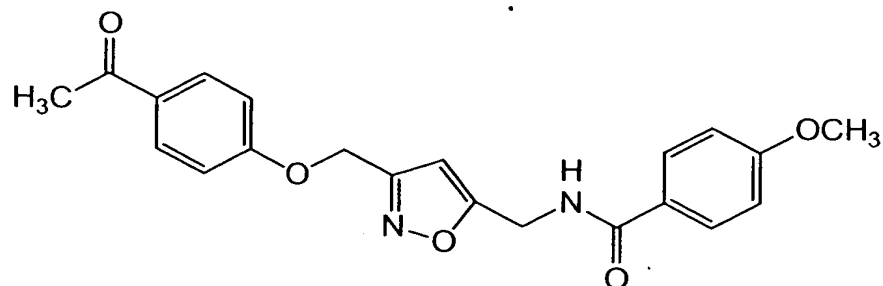


10

Melting point: 148 to 158°C.

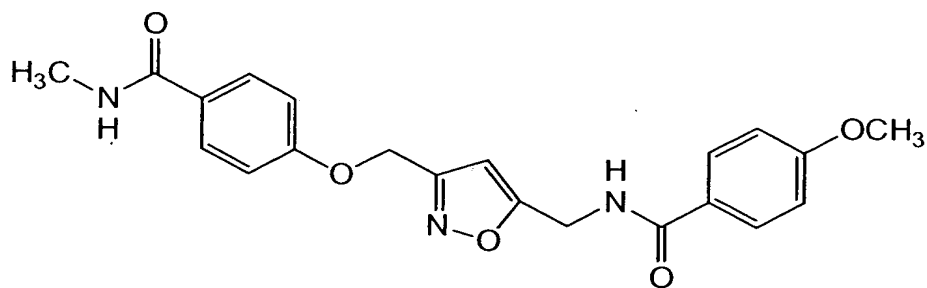
Example 18(135): Preparation of N-[[3-(4-acetylphenyl)oxymethyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-87):

15



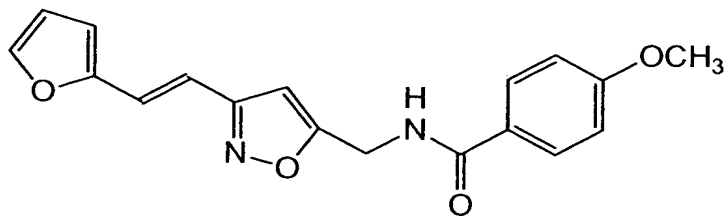
Melting point: 148 to 156°C.

Example 18(136): Preparation of N-{[3-(4-methylaminocarbonylphenyl)oxymethyl-5-isoxazolyl]methyl}-4-methoxybenzamide (Compound number IV-88):



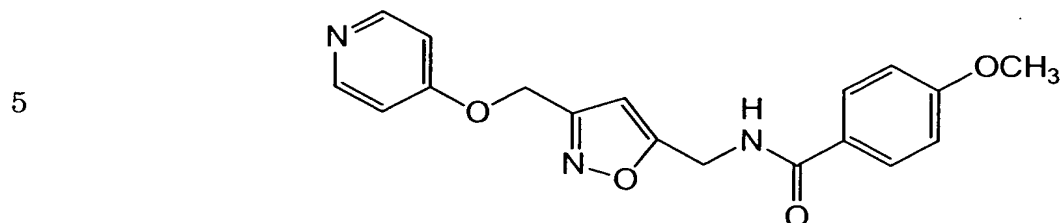
Melting point: 211 to 214°C.

10 Example 18(137): Preparation of N-{[3-(2-(2-furyl)ethenyl)-5-isoxazolyl]methyl}-4-methoxybenzamide (Compound number IV-89):



Melting point: 159 to 164°C.

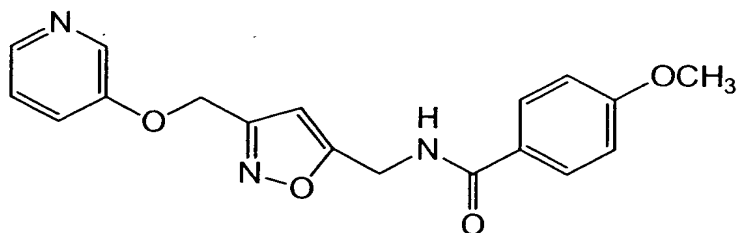
Example 18(138): Preparation of N-([3-(4-pyridyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-90):



Melting point: 154 to 159°C.

Example 18(139): Preparation of N-([3-(3-pyridyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-91):

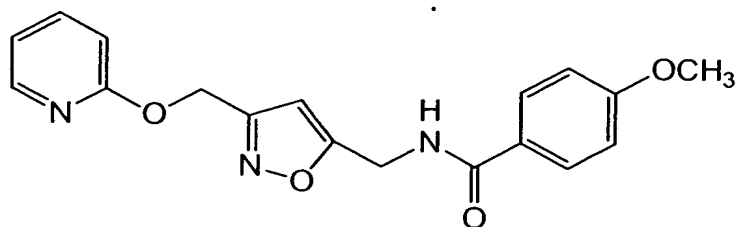
10



Property: oily;

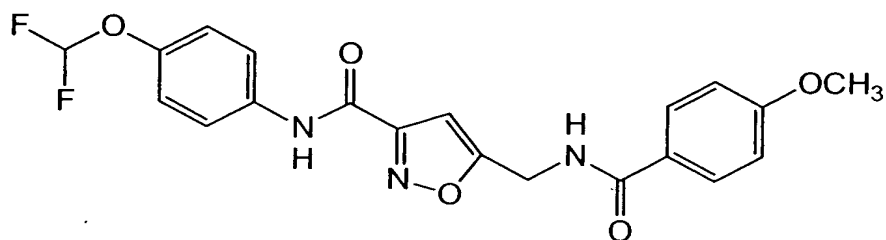
NMR :  $\delta$  3.78(3H, s), 4.66(2H, d), 5.08(2H, s), 6.32(1H, s), 6.70-8.35(9H, m)

15 Example 18(140): Preparation of N-([3-(2-pyridyl)oxymethyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-92):



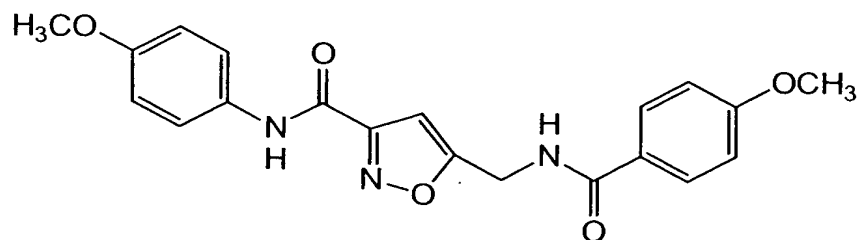
Melting point: 169 to 172°C.

Example 18(141): Preparation of N-([3-(4-difluoromethoxyphenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-93):



NMR :  $\delta$  3.85(3H, s), 4.76(2H, d), 6.60-8.00(9H, m), 8.30(1H, d), 9.30(1H, br)

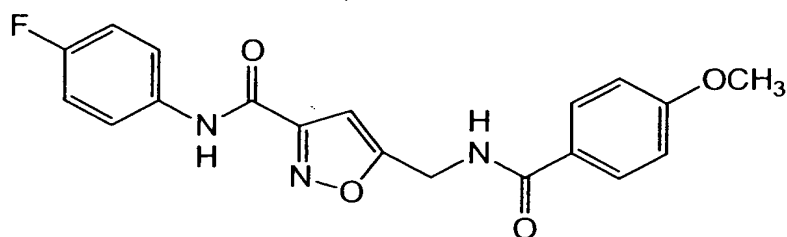
10 Example 18(142): Preparation of N-([3-(4-methoxyphenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-94):



Melting point: 186 to 188°C.

15 Example 18(143): Preparation of N-([3-(4-

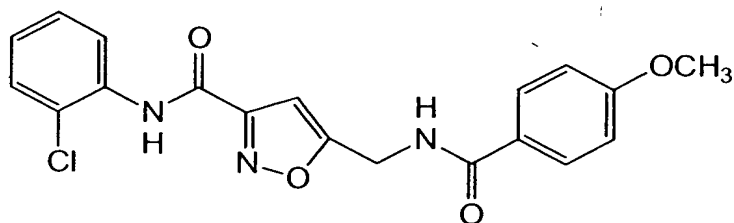
fluorophenyl)aminocarbonyl-5-isoxazolyl)methyl}-4-methoxybenzamide (Compound number IV-95):



Melting point: 191 to 193°C.

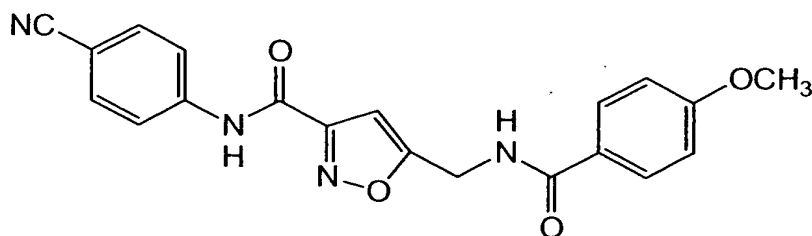
5

Example 18(144): Preparation of N-([3-(2-chlorophenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-96):



10 NMR :  $\delta$  3.83(3H, s), 4.76(2H, d), 6.60-8.00(9H, m), 8.20(1H, d), 9.10(1H, br)

Example 18(145): Preparation of N-([3-(4-cyanophenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-97):

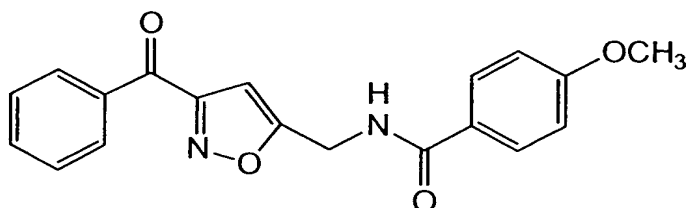


15

Melting point: 138 to 140°C.

Example 18(146): Preparation of N-[(3-benzoyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-98):

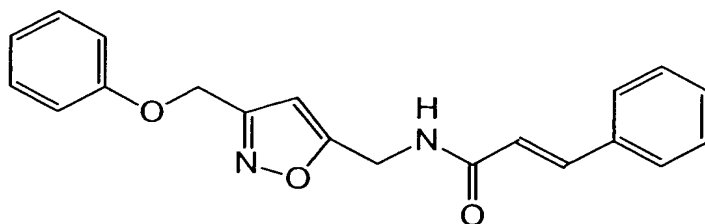
5



Melting point: 118 to 127°C.

Example 18(147): Preparation of N-[(3-phenoxyethyl-5-isoxazolyl)methyl]-β-phenylacrylamide (Compound number IV-99):

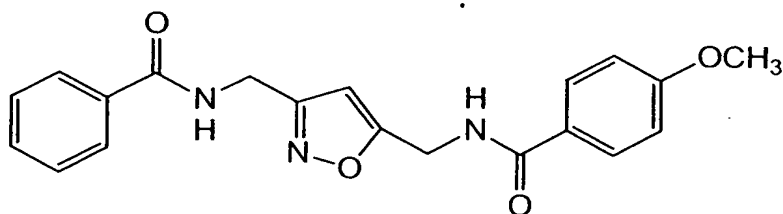
10



Melting point: 110 to 116°C.

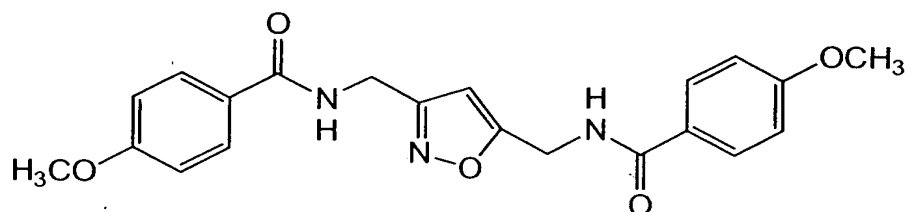
Example 18(148): Preparation of N-[(3-benzoylaminoethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-100):

15



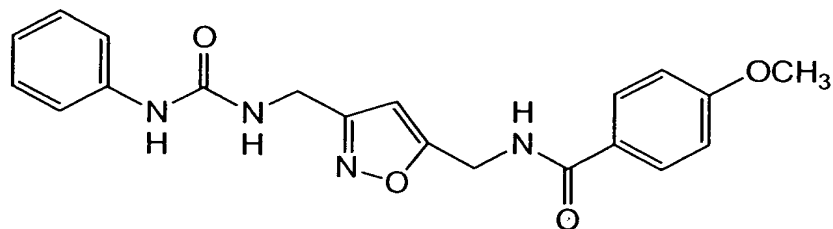
Melting point: 118 to 124°C.

Example 18(149): Preparation of N-[[3-(4-methoxybenzoyl)aminomethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-101):



Melting point: 176 to 179°C.

10 Example 18(150): Preparation of N-[(3-phenylaminocarbonylamino-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-102):



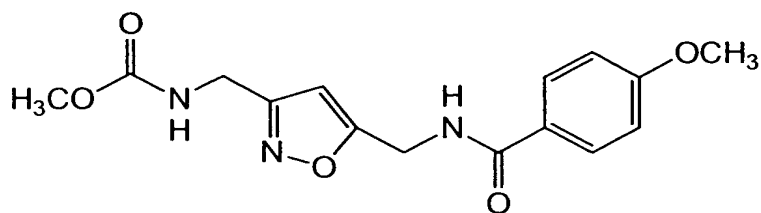
Melting point: 169 to 178°C.

15

Example 18(151): Preparation of N-[(3-



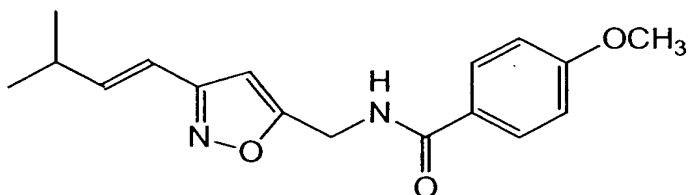
methoxycarbonylaminomethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-103):



Melting point: 143 to 145°C.

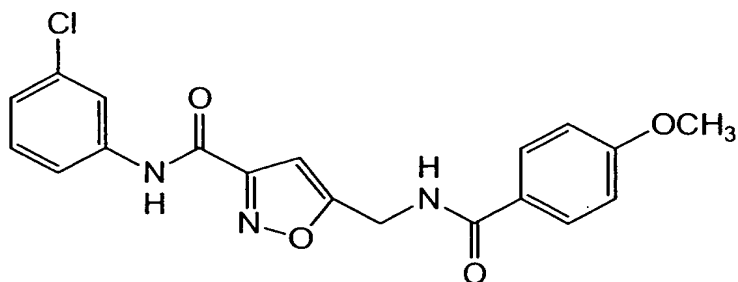
5

Example 18(152): Preparation of N-([3-(4-methyl-1-butenyl)-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-104):



10 Melting point: 104 to 113°C.

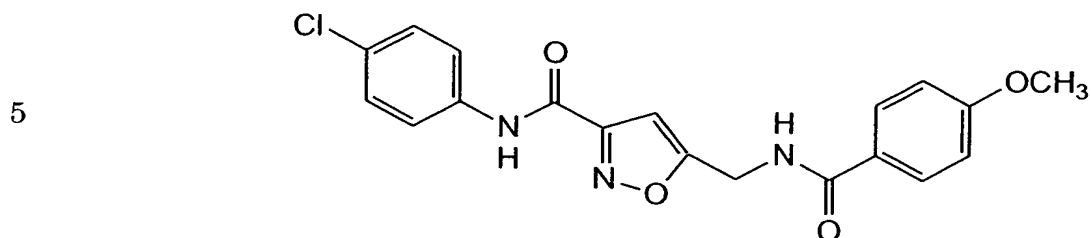
Example 18(153): Preparation of N-([3-(3-chlorophenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-105):



15

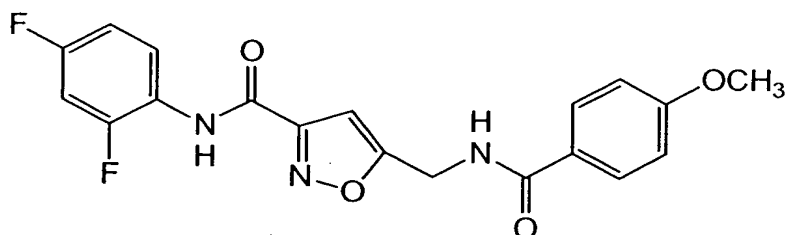
Melting point: 164 to 166°C.

Example 18(154): Preparation of N-([3-(4-chlorophenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-106):



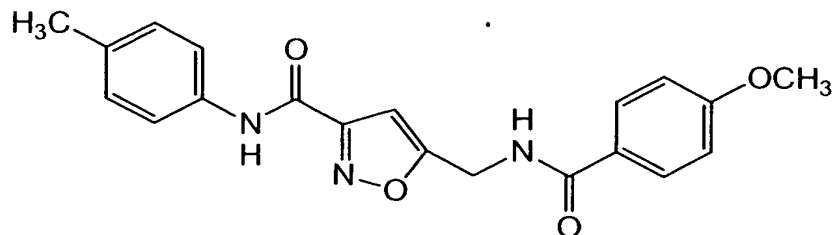
Melting point: 185 to 186°C.

Example 18(155): Preparation of N-([3-(2,4-difluorophenyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-107):



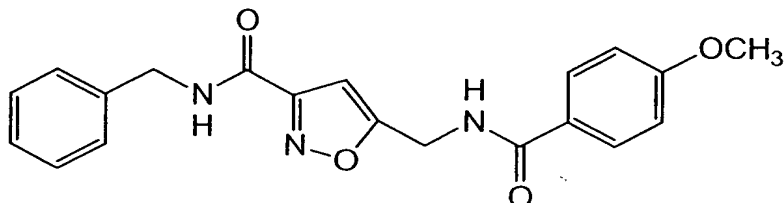
Melting point: 165 to 167°C.

Example 18(156): Preparation of N-([3-(4-tolyl)aminocarbonyl-5-isoxazolyl]methyl)-4-methoxybenzamide (Compound number IV-108):



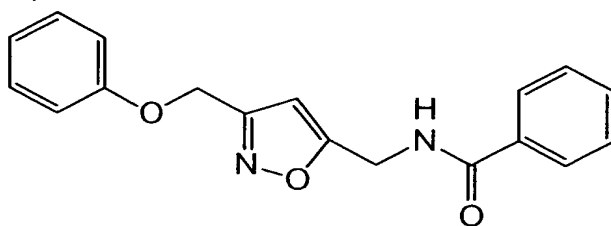
Melting point: 174 to 176°C.

Example 18(157): Preparation of N-[(3-benzylaminocarbonyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-109):



Melting point: 171 to 172°C.

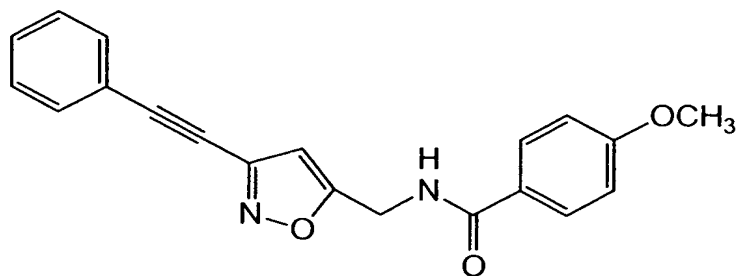
10 Example 18(158): Preparation of N-[(3-phenoxyethyl-5-isoxazolyl)methyl]benzamide (Compound number IV-44):



Melting point: 85 to 87°C.

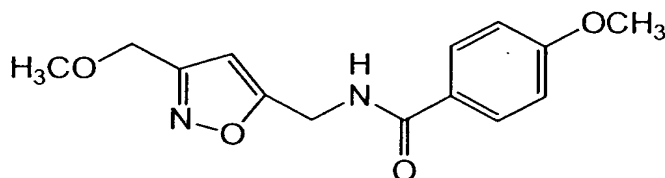
15 Example 18(159): Preparation of N-[(3-phenylethynyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-

110):



Melting point: 88 to 101°C.

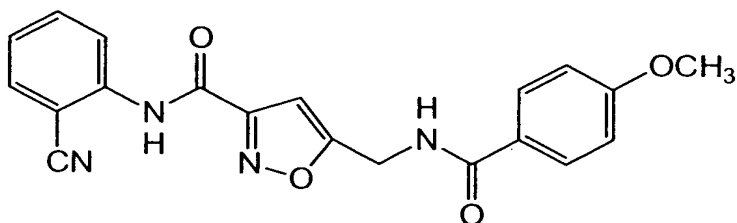
- 5 Example 18(160): Preparation of N-[(3-methoxymethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-111):



Melting point: 71 to 85°C.

10

- Example 18(161): Preparation of N-[[3-(2-cyanophenyl)aminocarbonyl-5-isoxazolyl]methyl]-4-methoxybenzamide (Compound number IV-112):

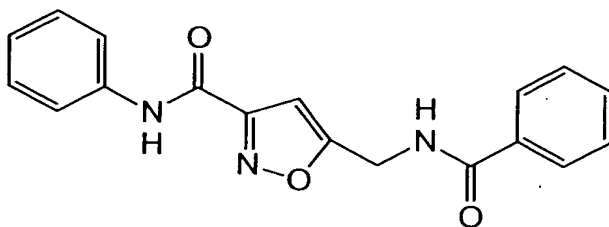


- 15 Property: gum-like;

NMR :  $\delta$  3.86(3H, s), 4.75(2H, d), 6.40-8.00(10H, m), 8.20(1H,

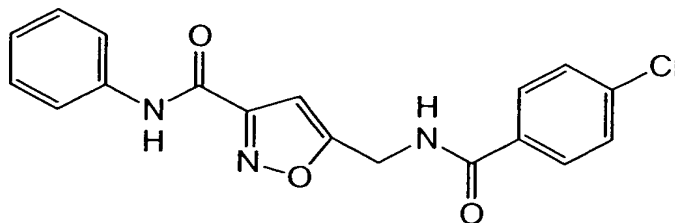
br), 9.40(1H, br)

Example 18(162): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]benzamide (Compound number IV-113):



5 Melting point: 177 to 178°C.

Example 18(163): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]-4-chlorobenzamide (Compound number IV-114):

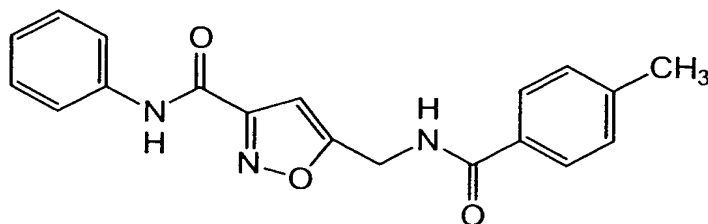


10

Melting point: 200°C or more.

Example 18(164): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]-4-methylbenzamide (Compound number IV-

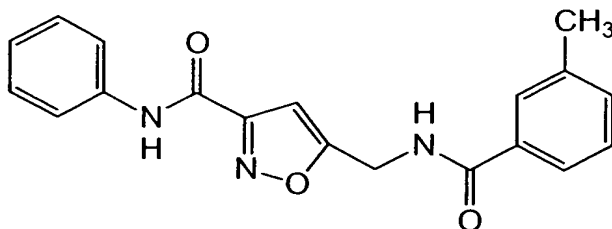
15 115):



Melting point: 193 to 195°C.

Example 18(165): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]-3-methylbenzamide (Compound number IV-

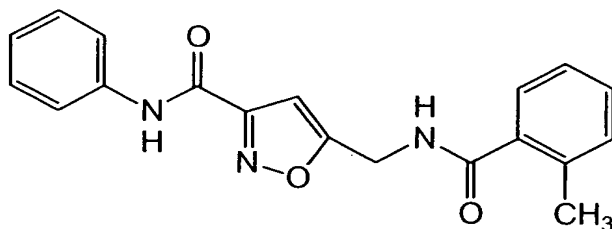
5 116):



Melting point: 151 to 152°C.

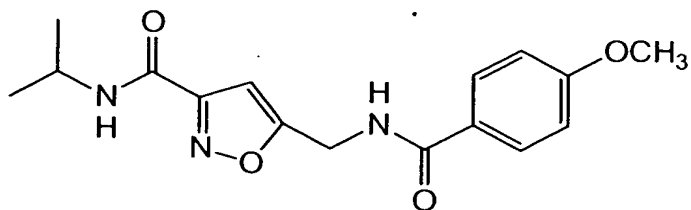
Example 18(166): Preparation of N-[(3-phenylaminocarbonyl-5-isoxazolyl)methyl]-2-methylbenzamide (Compound number IV-

10 117):



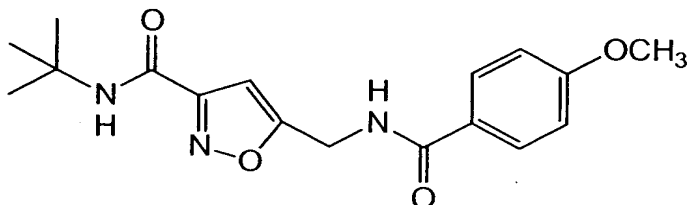
Melting point: 162 to 163°C.

15 Example 18(167): Preparation of N-[(3-isopropylaminocarbonyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-118):



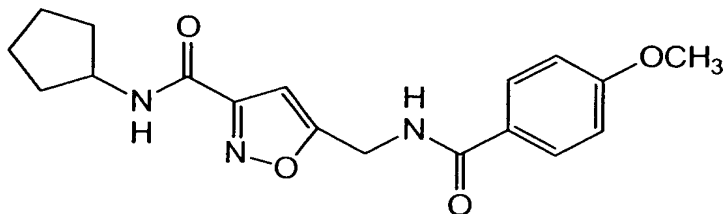
Melting point: 171 to 172°C.

Example 18(168): Preparation of N-[(3-tert-butylaminocarbonyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-119):



Melting point: 128 to 129°C.

10 Example 18(169): Preparation of N-[(3-cyclopentylaminocarbonyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-120):

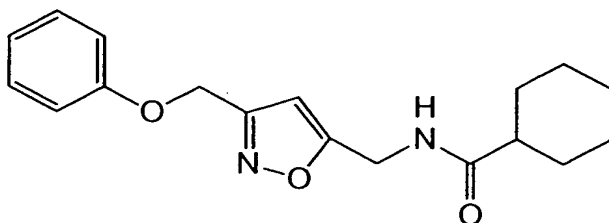


Melting point: 138 to 140°C.

15

Example 18(170): Preparation of N-[(3-phenoxyethyl-5-isoxazolyl)methyl]-4-methoxybenzamide

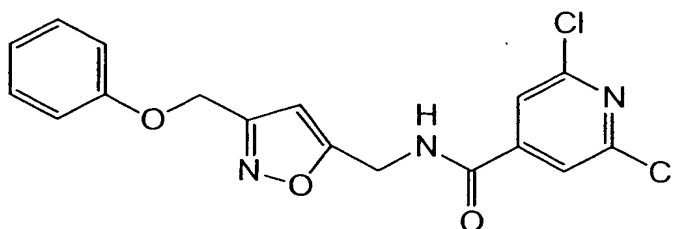
isoxazolyl)methyl]cyclohexanecarboxamide (Compound number IV-121):



Melting point: 86 to 93°C.

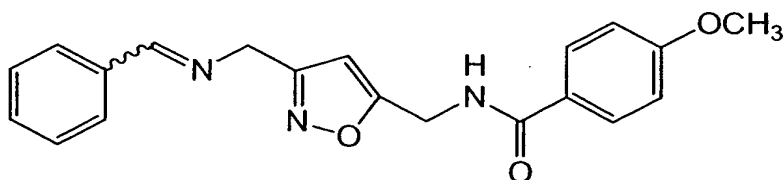
5

Example 18(171): Preparation of N-[(3-phenoxy-5-isoxazolyl)methyl]-2,6-dichloropyridine-4-carboxamide (Compound number IV-122):



10 Melting point: 93 to 99°C.

Example 18(172): Preparation of N-[(3-benzyliminomethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-123):



15

Property: amorphous;

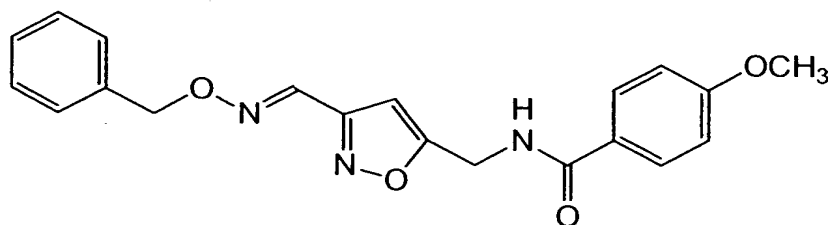
NMR :  $\delta$  3.74(3H, s), 4.66(2H, d), 4.74(2H, s), 6.23(1H, s),



6.84 (2H, d), 7.13-7.89 (7H, m), 8.35 (1H, s)

Example 18(173): Preparation of N-[(3-benzyloxyiminomethyl-5-isoxazolyl)methyl]-4-methoxybenzamide (Compound number IV-124):

5



NMR :  $\delta$  3.82 (3H, s), 4.71 (2H, d), 5.19 (2H, s), 6.51 (1H, s), 6.74 (1H, br), 6.89 (2H, d), 7.18-7.45 (5H, m), 7.75 (2H, d), 8.15 (1H, s)

#### 10 [Formulation Example]

##### Formulation Example 1

The compound of I-25 (16 parts by mass), 28 parts by mass of bentonite, 52 parts by mass of talc, 2 parts by mass of sodium dodecylbenzene sulfonate, and 2 parts by mass of sodium lignin sulfonate were mixed and milled. About 20 parts by mass of water was added, and the mixture was kneaded with a kneader, subsequently granulated through a granulator, then dried and sorted to afford granules.

#### 20 Formulation Example 2

The compound of II-3 (20 parts by mass), 30 parts by mass of diatomite, 36 parts by mass of clay, 5 parts by mass

of white carbon, 4 parts by mass of sodium lignin sulfonate, and 5 parts by mass of sodium alkyl naphthalene sulfonate were mixed and milled to afford water-dispersible powder.

#### 5 Formulation Example 3

The compound of III-12 (20 parts by mass), 65 parts by mass of kaolin clay, and 15 parts by mass of talc were thoroughly mixed and milled to afford 20% powder.

#### 10 Formulation Example 4

The compound of IV-8 (12.5 parts by mass), 4 parts by mass of ethylene glycol, 5 parts by mass of SORPOL 3940 (trade name, supplied from TOHO Chemical Industry Co., Ltd.), 1 part by mass of RUNOX 1000C (trade name, supplied from TOHO Chemical Industry Co., Ltd.), and 77 parts by mass of water were thoroughly mixed, and milled in a wet manner until an average particle size became 5  $\mu\text{m}$  or less. Subsequently, 0.5 part by mass of xanthane gum was mixed to afford a flowable agent.

20

#### Formulation Example 5

The compound of I-28 (20 parts by mass), 14 parts by mass of polyoxyethylene styrylphenyl ether, 10 parts by mass of calcium dodecylbenzene sulfonate, and 56 parts by mass of xylene were mixed to afford an emulsion.

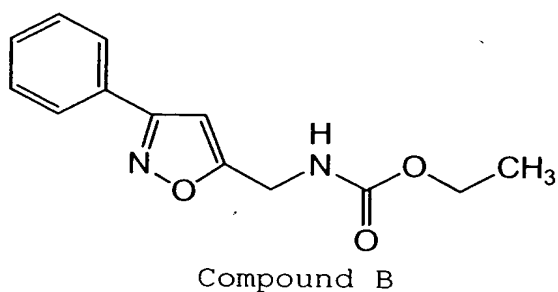
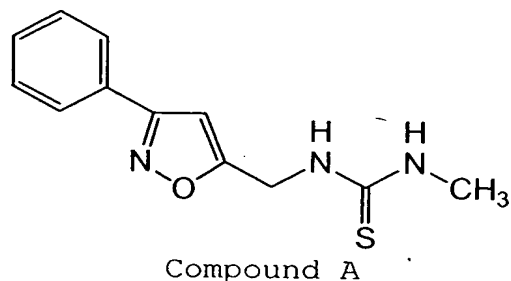
### Formulation Example 6

The compound of II-100 (17 parts by mass), 9.5 parts by mass of potassium chloride, 2 parts by mass of NEWCALGEN  
5 FS-1 (trade name, supplied from Takemoto Yushi Co., Ltd.), 1 part by mass of dextrin, and 0.5 part by mass of xanthane gum were mixed, and 20 parts by mass of water was added. The mixture was kneaded with a kneader, granulated through a granulator, and then dried/sorted to afford granules. Those  
10 granules were filled in a pouched container made from water-soluble film, and a filling opening was sealed with heat to afford a jumbo agent.

### [Evaluation Examples]

15 Next, Evaluation Examples of the compounds of the invention as the agri-horticultural fungicides are described.

The following compounds A and B described in JP 52-154525 A were used as reference drugs.



20

### Evaluation Example 1. Effect test for rice blast

The formulation of the compound of the invention prepared according to Formulation Example 2 was diluted with water to have a concentration of 200 ppm, and subsequently a sufficient amount of the solution was sprayed with a spraygun on rice (cultivar: Koshihikari, planting 15 seedlings) in a 3 leaf stage grown in a plastic pot with a diameter of 6 cm. Next day, a sporal suspension of *Pyricularia oryzae* which was a causative fungus of the rice blast was sprayed to inoculate. The pot was retained at 22°C under a wet condition for 24 hours, subsequently left in a green house for 7 days. Then, lesion numbers on the third leaves were grossly surveyed, and evaluated according to the criteria shown in Table 5. The results are shown in Table 6.

Table 5

Evaluation	Lesion area rate (%)
5	0
4	1~5
3	6~10
2	11~20
1	21~41
0	41 or more

Table 6-1: Effect for rice blast

Compound No.	Example No.	Evaluation	Compound No.	Example No	Evaluation
I-4	17(1)	5	II-106	18(12)	3
I-6	17(2)	5	II-108	18(14)	1
I-8	18(3)	3	II-110	18(16)	1
I-25	17(3)	5	II-112	18(18)	3
I-26	5	5	II-113	18(19)	2
I-27	17(4)	5	II-114	18(20)	3
I-28	1	4	II-115	18(21)	3
I-29	17(5)	5	III-2	17(29)	4
I-30	17(6)	5	III-4	17(30)	3
I-31	12	4	III-5	9	4
I-32	17(7)	5	III-6	17(31)	5
I-33	17(8)	5	III-10	17(32)	5
I-38	17(9)	5	III-11	17(33)	5
I-41	17(10)	3	III-12	17(34)	5
I-45	17(12)	4	III-13	17(35)	5
I-47	17(14)	4	III-14	17(36)	5
I-48	17(15)	4	III-17	17(37)	5
I-49	17(16)	3	III-18	7	5
I-63	18(2)	2	III-19	17(38)	5
I-64	18(4)	2	III-26	17(39)	5
II-2	6	5	III-27	17(40)	5
II-3	17(17)	3	III-28	17(41)	5
II-7	11	5	III-31	17(42)	5
II-9	17(19)	5	III-34	17(43)	5
II-10	18(9)	4	III-37	17(44)	5
II-12	3	4	III-40	17(45)	5
II-13	17(20)	5	III-43	17(46)	5
II-14	17(21)	4	III-49	17(47)	5
II-15	17(22)	3	III-51	17(48)	5
II-48	17(23)	3	III-52	17(49)	5
II-49	17(24)	4	III-57	17(51)	2
II-50	17(25)	5	III-61	17(52)	5
II-99	17(26)	3	III-65	17(53)	5
II-100	17(27)	4	III-66	17(54)	5
II-104	18(10)	5	III-67	17(55)	5
II-105	18(11)	2	III-68	17(56)	5

Table 6-2 : Effect for rice blast

Compound No.	Example No.	Evaluation	Compound No.	Example No.	Evaluation
III-73	17(57)	5	III-223	18(38)	5
III-74	17(58)	5	III-224	18(42)	2
III-107	17(59)	5	III-225	18(43)	4
III-108	17(60)	2	III-227	18(45)	5
III-109	17(61)	4	III-228	18(46)	4
III-111	4	5	III-229	18(47)	5
III-121	17(62)	5	III-230	18(48)	5
III-122	17(63)	4	III-231	18(49)	5
III-123	17(64)	5	III-232	18(50)	3
III-124	17(65)	5	III-235	18(53)	4
III-130	17(66)	4	III-236	18(54)	4
III-133	17(67)	4	III-237	18(55)	3
III-134	17(68)	3	III-238	18(56)	4
III-137	17(69)	4	III-240	18(59)	2
III-139	17(70)	5	III-243	18(62)	2
III-156	17(71)	4	III-245	18(64)	5
III-162	17(72)	5	III-246	18(65)	3
III-174	17(73)	4	III-247	18(66)	5
III-177	17(76)	4	III-248	18(67)	3
III-180	17(77)	5	III-249	18(68)	2
III-201	17(78)	5	III-250	18(69)	2
III-203	18(40)	2	III-251	18(70)	5
III-204	18(37)	2	III-252	18(71)	5
III-208	17(80)	4	III-253	18(72)	5
III-210	17(82)	5	III-255	18(74)	4
III-212	18(26)	2	III-256	18(75)	5
III-213	18(27)	3	III-257	18(76)	3
III-214	18(28)	4	III-258	18(77)	5
III-215	18(29)	5	III-259	18(78)	5
III-216	18(30)	4	III-262	18(81)	5
III-217	18(31)	4	III-263	18(82)	5
III-218	18(32)	3	III-264	18(83)	5
III-219	18(33)	3	III-265	18(84)	5
III-220	18(34)	3	III-275	18(94)	2
III-221	18(35)	2	IV-1	17(84)	5
III-222	18(36)	3	IV-2	17(85)	5

Table 6-3:Effect for rice blast

Compound No.	Example No.	Evaluation	Compound No.	Example No.	Evaluation
IV-3	17(86)	5	IV-84	18(131)	5
IV-4	17(87)	4	IV-86	18(133)	5
IV-5	17(88)	4	IV-87	18(135)	5
IV-6	17(89)	5	IV-88	18(136)	3
IV-8	17(91)	3	IV-89	18(137)	5
IV-9	17(92)	5	IV-90	18(138)	3
IV-10	17(93)	5	IV-91	18(139)	3
IV-11	17(94)	5	IV-92	18(140)	4
IV-12	8	5	IV-93	18(141)	4
IV-29	17(96)	3	IV-94	18(142)	2
IV-33	17(97)	5	IV-95	18(143)	4
IV-34	17(98)	5	IV-96	18(144)	4
IV-35	17(99)	5	IV-97	18(145)	5
IV-40	17(100)	3	IV-98	18(146)	2
IV-44	18(158)	5	IV-99	18(147)	4
IV-45	10	5	IV-104	18(152)	4
IV-46	17(101)	5	IV-105	18(153)	3
IV-47	16	5	IV-106	18(154)	2
IV-48	17(102)	3	IV-107	18(155)	5
IV-49	17(103)	2	IV-108	18(156)	2
IV-50	17(104)	5	IV-110	18(159)	5
IV-51	17(105)	2	IV-111	18(160)	3
IV-52	17(106)	3	IV-112	18(161)	2
IV-53	17(107)	5	IV-113	18(162)	4
IV-54	17(108)	3	IV-114	18(163)	5
IV-55	17(109)	4	IV-115	18(164)	5
IV-56	17(110)	3	IV-116	18(165)	5
IV-57	17(111)	5	IV-117	18(166)	4
IV-58	17(112)	5	IV-119	18(168)	2
IV-59	17(113)	5	IV-120	18(169)	3
IV-60	17(114)	5	IV-121	18(170)	4
IV-72	17(115)	2	IV-122	18(171)	3
IV-73	18(119)	2	IV-123	18(172)	2
IV-74	17(116)	4	IV-124	18(173)	5
IV-75	18(120)	2	Compound A		0
IV-80	18(127)	4	Compound B		0

### Evaluation Example 2. Effect test for cucumber anthracnose

The formulation of the compound of the invention prepared according to Formulation Example 2 was diluted with water to have a concentration of 200 ppm, and subsequently a  
5 sufficient amount of the solution was sprayed with a spraygun on the first and second leaves of cucumber (cultivar: Tokiwa Hikari 3-go P-type) in a 3 leaf stage grown in a plastic pot with a diameter of 6 cm. Next day, a sporal suspension of *Colletotrichum lagenarium* which was a  
10 causative fungus of the cucumber anthracnose was sprayed to inoculate. The pot was retained at 22°C under a wet condition for 24 hours, subsequently left in a green house for 7 days. Then, lesion numbers on the first and second leaves were grossly surveyed, and evaluated according to the  
15 same criteria as those shown in Table 5. The results are shown in Table 7.



Table 7-1: Effect for cucumber anthracnose

Compound No.	Example No.	Evaluation	Compound No.	Example No.	Evaluation
I-4	17(1)	5	III-11	17(33)	3
I-6	17(2)	5	III-12	17(34)	5
I-8	18(3)	3	III-13	17(35)	5
I-26	5	5	III-14	17(36)	3
I-27	17(4)	5	III-17	17(37)	5
I-28	1	5	III-18	7	5
I-29	17(5)	5	III-19	17(38)	5
I-30	17(6)	5	III-26	17(39)	5
I-33	17(8)	5	III-27	17(40)	5
I-38	17(9)	5	III-28	17(41)	4
I-41	17(10)	4	III-31	17(42)	5
I-45	17(12)	4	III-34	17(43)	5
I-47	17(14)	4	III-37	17(44)	3
I-64	18(4)	3	III-40	17(45)	5
II-7	11	5	III-43	17(46)	4
II-9	17(19)	5	III-49	17(47)	5
II-10	18(9)	5	III-51	17(48)	4
II-12	3	4	III-52	17(49)	5
II-13	17(20)	5	III-61	17(52)	4
II-14	17(21)	5	III-65	17(53)	5
II-15	17(22)	4	III-66	17(54)	5
II-48	17(23)	4	III-67	17(55)	5
II-100	17(27)	4	III-68	17(56)	5
II-104	18(10)	5	III-73	17(57)	4
II-106	18(12)	2	III-74	17(58)	5
II-108	18(14)	2	III-107	17(59)	5
II-110	18(16)	1	III-108	17(60)	3
II-112	18(18)	3	III-109	17(61)	5
II-113	18(19)	1	III-111	4	5
II-114	18(20)	3	III-121	17(62)	5
II-115	18(21)	2	III-122	17(63)	4
III-2	17(29)	4	III-123	17(64)	5
III-4	17(30)	4	III-124	17(65)	4
III-5	9	4	III-130	17(66)	4
III-6	17(31)	4	III-133	17(67)	4
III-10	17(32)	5	III-134	17(68)	5

Table 7-2: Effect for cucumber anthracnose

Compound No.	Example No.	Evaluation	Compound No.	Example No.	Evaluation
III-137	17(69)	5	III-246	18(65)	3
III-139	17(70)	5	III-247	18(66)	5
III-156	17(71)	4	III-248	18(67)	2
III-162	17(72)	5	III-249	18(68)	2
III-174	17(73)	4	III-250	18(69)	2
III-177	17(76)	4	III-251	18(70)	5
III-180	17(77)	5	III-252	18(71)	5
III-201	17(78)	5	III-253	18(72)	5
III-204	18(37)	2	III-255	18(74)	4
III-208	17(80)	4	III-256	18(75)	5
III-210	17(82)	5	III-257	18(76)	4
III-214	18(28)	5	III-258	18(77)	4
III-215	18(29)	5	III-259	18(78)	4
III-216	18(30)	4	III-262	18(81)	5
III-217	18(31)	4	III-263	18(82)	5
III-218	18(32)	4	III-264	18(83)	5
III-219	18(33)	3	III-265	18(84)	4
III-220	18(34)	3	III-275	18(94)	2
III-221	18(35)	2	IV-1	17(84)	5
III-222	18(36)	2	IV-2	17(85)	5
III-223	18(38)	5	IV-3	17(86)	5
III-224	18(42)	2	IV-4	17(87)	4
III-225	18(43)	4	IV-9	17(92)	5
III-227	18(45)	5	IV-10	17(93)	5
III-228	18(46)	3	IV-11	17(94)	5
III-229	18(47)	5	IV-12	8	5
III-230	18(48)	5	IV-29	17(96)	3
III-231	18(49)	5	IV-33	17(97)	5
III-232	18(50)	2	IV-34	17(98)	5
III-235	18(53)	4	IV-35	17(99)	5
III-236	18(54)	4	IV-40	17(100)	2
III-237	18(55)	3	IV-44	18(158)	5
III-238	18(56)	5	IV-45	10	5
III-240	18(59)	2	IV-46	17(101)	5
III-243	18(62)	1	IV-47	16	5
III-245	18(64)	5	IV-48	17(102)	2

Table 7-3:Effect for cucumber anthracnose

Compound No.	Example No.	Evaluation	Compound No.	Example No.	Evaluation
IV-49	17(103)	2	IV-95	18(143)	4
IV-50	17(104)	4	IV-96	18(144)	4
IV-52	17(106)	2	IV-97	18(145)	5
IV-53	17(107)	5	IV-99	18(147)	4
IV-54	17(108)	4	IV-104	18(152)	4
IV-55	17(109)	4	IV-105	18(153)	3
IV-56	17(110)	2	IV-107	18(155)	5
IV-57	17(111)	5	IV-108	18(156)	2
IV-58	17(112)	5	IV-110	18(159)	5
IV-59	17(113)	5	IV-111	18(160)	3
IV-60	17(114)	5	IV-112	18(161)	3
IV-72	17(115)	3	IV-113	18(162)	4
IV-73	18(119)	3	IV-114	18(163)	5
IV-74	17(116)	3	IV-115	18(164)	5
IV-75	18(120)	3	IV-116	18(165)	4
IV-80	18(127)	4	IV-117	18(166)	4
IV-84	18(131)	5	IV-119	18(168)	3
IV-86	18(133)	5	IV-120	18(169)	2
IV-87	18(135)	4	IV-121	18(170)	4
IV-88	18(136)	3	IV-122	18(171)	2
IV-89	18(137)	5	IV-123	18(172)	2
IV-90	18(138)	3	IV-124	18(173)	5
IV-92	18(140)	3	Compound A		0
IV-93	18(141)	3	Compound B		0
IV-94	18(142)	3			

Evaluation Example 3. Effect test for tomato late blight

The active ingredient compound was diluted with water  
5 containing 0.01% Tween 20 to have a concentration of 125 ppm.  
The amount of this drug solution corresponding to 200 L/10 a  
was sprayed with a spraygun on tomato in a 5 leaf stage  
(cultivar: Sugar Lump) grown in a plastic pot with a

diameter of 6 cm. Next day, a zoosporangium suspension of *Phytophthora infestans* which was the causative fungus of tomato late blight was sprayed to inoculate. The pot was retained at 22°C under a wet condition for 16 hours, then  
5 left in a green house for 3 days. Subsequently, diseased area rates of the third, fourth and fifth true leaves were grossly surveyed, and evaluated according to the same criteria as those in Table 5. The results are shown in Table 8.

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Table 8:Effect for tomato late blight

Compound No.	Example No.	Evaluation
I -65	18 (5)	4
I -67	18 (8)	2
II -105	18 (11)	5
II -107	18 (13)	4
II -111	18 (17)	5
II -115	18 (21)	3
III -2	17 (29)	5
III -3	15	2
III -226	18 (44)	3
III -233	18 (51)	2
III -254	18 (73)	5
III -260	18 (79)	3
III -261	18 (80)	2
III -266	18 (85)	1
III -267	18 (86)	5
III -268	18 (87)	0
III -269	18 (88)	5
III -270	18 (89)	2
III -271	18 (90)	1
III -272	18 (91)	5
III -273	18 (92)	0
III -274	18 (93)	4
III -277	18 (96)	4
III -278	18 (97)	4
III -279	18 (98)	5
IV -78	18 (125)	1
Compound A		0
Compound B		0

Evaluation Example 4. Effect test for cucumber downy mildew

The active ingredient compound was diluted with water containing 0.01% Tween 20 to have a concentration of 125 ppm.

- 5 The amount of this drug solution corresponding to 200 L/10 a was sprayed with a spraygun on cucumber in a 3 leaf stage (cultivar: Hikari 3-go P-type) grown in a plastic pot with a

diameter of 6 cm. Next day, a zoosporangium suspension of *Pseudoperonospora cubensis* which was the causative fungus of cucumber downy mildew was sprayed to inoculate. The pot was retained at 22°C under a wet condition for 18 hours, then left in a green house for 3 days. Subsequently, diseased area rates of the first and second true leaves were grossly surveyed, and evaluated according to the same criteria as those in Table 5. The results are shown in Table 9.

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Table 9: Effect for cucumber downy mildew

Compound No.	Example No.	Evaluation
I-65	18(5)	2
II-105	18(11)	4
II-107	18(13)	2
II-111	18(17)	3
II-115	18(21)	1
III-2	17(29)	5
III-3	15	4
III-204	18(37)	4
III-222	18(36)	4
III-226	18(44)	4
III-254	18(73)	3
III-267	18(86)	5
III-269	18(88)	5
III-272	18(91)	3
III-274	18(93)	2
III-277	18(96)	2
III-278	18(97)	2
III-279	18(98)	3
Compound A		0
Compound B		0

#### INDUSTRIAL APPLICABILITY

The substituted isoxazole alkylamine derivative represented by the formula (I) or (IA) exhibits the disease  
5 control effect on plant disease damages, particularly on rice blast and cucumber anthracnose, in a low drug amount, is safe for crops, and is useful as the agri-horticultural fungicide.